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# FOREIGN TECHNOLOGY DIVISION



## ALL UNION CONGRESS ON CHEMISTRY AND THE APPLICATION OF ADDITIVES TO LUBRICANTS AND FUELS

(Selected Articles)

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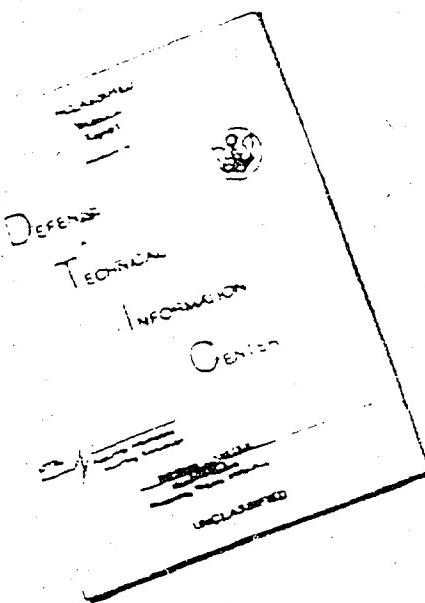


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## EDITED TRANSLATION

ALL UNION CONGRESS ON CHEMISTRY AND THE APPLICATION OF ADDITIVES  
TO LUBRICANTS AND FUELS

(Selected Articles)

English pages: 122

SOURCE: Vsesoyuznoye Soveshchaniye po Khimii i Primeneniyu  
Prisadok K Maslam i Toplivam 2D, 1966; Prisadok K  
Maslam, Trudy (All Union Congress on Chemistry and  
the Application of Additives to Lubricants and Fuels).  
2nd Report, 1966 Additives to Lubricants. Transactions.  
1966, pp. 148-157, 281-324, 336-341, 370-388.

Translated Under: Contract #33657-68-D-0866 P002

THIS TRANSLATION IS A REPORT ON THE SCIENTIFIC,  
TECHNICAL, AND PRACTICAL STATEMENTS ON THE  
APPLICATION OF ADDITIVES TO LUBRICANTS AND FUELS.  
IT REFLECTS THE PRACTICAL AND THEORETICAL  
ACHIEVEMENTS OF THE RUSSIAN SCIENCE AND  
TECHNOLOGY IN THIS FIELD.

FID - HT - 23-1496-68

Date: 1 June 1969

DATA HANDLING PAGE	
SEARCHED	SEARCHED LOC.
TF-900097	20-SECURITY CLASS
07, 11	SECRET
01-SECURITY AND CONFIDENTIAL INFORMATION	SECRET
UNCL. D	UNCL.
02-AUTHORS KREYN, S. E.; SHKRETER, YU. N.; TITOV, LEVITIN M. K.; 16-VILENNIK, A. V.; 16-KALASHNIKOV, V. P.	02-DOCUMENT NO. HT-25-1196-68 03-PROJECT NO. 73201-78
03-DATE OF PUBLICATION 1966 VSFSOZNUYE SOVESHCHANIYE PO KHIMII I PRIMENENIYU PRISADOK K MASLAM I TOPLIVAN 2D, 1966, PRISADOK K MASLAM. TRUDY (RUSSIAN)	04-CONTROLS, BARRIERS IN MEASURE CLASSE
05-REEL FRAME NO. 1089 0532	06-CHANGES 70-CHANGES
07-REP. ACC. NO. F33657-68-D- 0886 R002	08-REP. DATE 54-00
08-STEP NO. 02-UR/0000/66/000/000/0148/0157	09-TRANSLATION NONE
ABSTRACT	
(U) The purpose of this investigation was to improve the antifriction and corrosion inhibiting properties of automotive lubricating oils. The experiments involved nitration of mineral oils. On the basis of these experiments an oil additive AKOR-1, and technical production methods for this additive were developed. AKOR-1 improves the corrosion inhibiting properties of motor and transmission oils with respect to cast iron and nonferrous metals. AKOR-1 improves only the anticorrosive and cleaning properties of oils, while the other properties of the oils remain the same. AKOR-1 added in 10 percent concentration to motor and lubricating oil successfully passed practical field tests in the internal combustion engines OD-9, 2Ch-8, 5A1, GAZ-69, and YaAZ-238 and in trucks GAZ-51, ZiL-157, and KRAZ-214. ORIG. art. has: 2 formulas, 2 figures, and 6 tables.	

DATA NAMEPLATE PAGE		SECURITY TAB	
ST-ACCESN. NO. 2-ACCESN. LOC.	11, 13	AS-8 OPERATING EXPERIENCE WITH ZIL-130 ENGINES USING AS-8 OIL WITH VNIIT KP-360 ADDITIVE	VNIIT KISELEVKA, T. T. - 16-BURSEVSKA, S. V.
AS-8 SUBJECT AREA		AS-8 DATE OF INFO 7/1/63 AS-8 DOCUMENT NO. FPP-12-1400-63 AS-8 PROJ. NO. 72-001-72 AS-8 TESTS CLOSING	
AS-8 AUTHOR CO-AUTHORS OBLEUKHOVA, O. S. I. FROTOSON, V. V. TIE- KISELEVKA, T. T. - 16-BURSEVSKA, S. V.		UNCL	
AS-8 SOURCE VSEGOZDNOYE SOVESHCHANIYE PO KHIMII I PRIMENENIYU PRISADOK K MASLAM I TOPELIYU 2D, 1966, PRISADKI K MASLAM. TRUDY (RUSSIAN)		AS-8 PAGES 7 REVISED PAGE	
AS-8 SECURITY AND INFORMATION AND INFORMATION		AS-8 COMMENTS NONE	
UNCL. O			
AS-8 FILE NUMBER 77-3098888		AS-8 CHANGES UP	
1889. 05.31 CONTRACT NO. F53657-65-D- 0886. R002		AS-8 DATE PROBLEMS DATE 54-00 TRANSMISSION ACCESSION NO. 02-JR/0000/66/000/000/0281/0286	
ABSTRACT			
(V) The performance of the LIAZ rpm. ZIL-130 engine, running on AS-8 lubricating oil with VNIIT KP-360 additive was tested in a fleet of 70 motor vehicles operating under various climatic conditions. The purpose of the study was to examine the lubricating properties of an oil prepared according to standard GOST 10541-63. The study involved visual examination of the engine parts in order to assess the detergent-, anticorrosive-, antiscuffing-, and low temperature properties of the lubricating oil. In addition, micrometric measurements were made to determine the oil's anti-wear property and the chemical analysis of the used oil was employed for determining the oil deterioration. The tests (200,000 kilometers) revealed that the AS-8 oil with VNIIT KP-360 additive increases the wear resistance of the parts of the ZIL-130 engine. Orig. art. han. figure 3 tables.			

DATA HANDLING PAGE	
16-ACCNUM OR NO.	16-ACCNUM OR NO.
T90000909	16-TITLE TEST RESULTS FOR OILS WITH ADDITIVES USED IN THE YAMZ-238NB ENGINE
11, 13	11-PROJECT AREA
MANUFACT CO-AUTOMOBIL SUGAR, N. I. IIG-SJARON, YE. P.; 16-YEFREMOV, V. N.; 16-GUSEV, G. M.; 16-BELYACHIKOV, G. P.	16-STATE OF INFO -66 16-DOCUMENT NO. FD-11-25-1166-66 16-PROJ. NO. 72301-78
16-TESTS OF UZNOYE SOVESHCHANIYE PA KHIMII I PRIMENENIYA PRISADOK K MASTIAM I TOPLIVAM 2D. 1966, PRISADOK K MASTIAM. TRUDY (RUSSIAN)	16-HEADER CL. A&B
16-SECURITY AND CONFIDENTIALITY INFORMATION UNCL. O	16-CONTROL MARKINGS UNCL.
16-REFL. PAGE NO. 17-REFL. PAGE NO.	16-GEOPHICAL AREA UR
1889 0534	17-CHANCES TYPE PRODUCT TRANSLATION ACCESSION NO.
CONTRACT NO. F73657-68-D-0866 P002 FSP NO. 02-UR/0000/66/000/000/0286/0290	17-PUBLISHING DATE 94-00 NONE 7
ABSTRACT	
(U) The performance of several batches of lubricating oils with various additives were tested in a new four-stroke diesel engine (Yamz-238NB) developed by the Yaroslav Engine Plant (Yaroslavskiy motorny zavod). The oils were evaluated on a standardized stationary engine run at different rpm's (1500-1700) for various periods (10-110 minutes) under different loads. The results of the tests were found to be very inconclusive because of the variation in the properties of the base oil in various batches. However, the best performance in the Yamz-238NB engine was exhibited by Series 1 oil supplemented with 5 percent VNIIT NP-370, 2 percent PMS-200A, and 0.005 percent E2-5K, and 0.005 percent PMS-200A additives. Orig. art. has: 6 figures, 3 tables.	

(U) The performance of several batches of lubricating oils with various additives were tested in a new four-stroke diesel engine (YAMZ-238MB) developed by the Yaroslavl Engine Plant (Yaroslavskiy motornyy zavod). The oils were evaluated on a standardized stationary engine run at different rpm's (1,500-1,700) for various periods (10-20 minutes) under different loads. The results of the tests were found to be very inconclusive because of the variation in the properties of the base oil in various batches. However, the best performance in the YAMZ-238MB engine was exhibited by Series 1 oil supplemented 5 percent VNIIT NP-370, 2 percent PIC Ya, 0.5 percent 2-25K, and 0.05 percent PMS-2004 additives. Orig. art. has 24 tabs.

(V) Several additives (MASE, DF-111, PES-2004) were examined after improvement of the characteristics of VRII NP-1 transmission oil for passenger cars. The tests were based on visual examination of the different parts of antifriction operating after various distances (up to 100,000 kilometers) under various climatic conditions and ambient temperatures (minus 60 degrees to plus 130 degrees C.). The best oil performance (2.2 percent longer service) resulted after supplementing the VRII NP-1 transmission oil base with 4 percent of MASE, 2.2 percent of DF-111, and 0.005 percent of PES-2004 additives. The present marked transmission oil (VRII NP-1) contains DF-111 additive and its antigum, antiwear, antioxidant, and anticorrosion properties are considered unsatisfactory.

DATA SHEET PAGE		SUBJECT AREA	
PT5000911	20. Topic Title lubricant additive, lubricant property, lubricating oil, diesel engine	1. Accession no.	2. Report No.
a. INTERCOMBINATION OF ASH-FORMING AND POLYMERIC ASH-FREE ADDITIVES IN MOTOR OILS AS A WAY TO FURTHER QUALITY IMPROVEMENT		3. Date of info	4. Date of info
		1989-07-16	1989-07-16
		5. SOURCE	6. INVENTOR
b. VSEGOVNOYE SODESHCHENIYE PO KHMILI I PRIMENENIYU PRISADOK K MASLAM I TOPIL'YU (RUSSIAN)		7. DOCUMENT NO.	8. INVENTOR NO.
2D, 1986, PRISADOK K MASLAM. TRUDY (RUSSIAN)		PTD-ET-23-149-12	FTD-ET-23-149-6
c. SECURITY AND DOWNGRADING INFORMATION		9. SECURITY CLASSIFICATION	10. COUNTRY, ORIGINATOR
		UNCL	UNCL
d. COUNTRY, NATION		11. COUNTRY	12. SECURITY CLASSIFICATION
		UR	UNCL
e. PUBLICATIONS		13. PUBLICATION DATE	14. PUBLICATION DATE
f. NEW ADD. NO.		15. TYPE PROPERTY	16. TYPE PROPERTY
P-33657-68-D-0856-P002		91-00 TRANSLATION	91-00 TRANSLATION
g. STEP NO.		17. ACCESSION NO.	18. SUBJECT
02-UR/0000/66/000/0297/0500		02-UR/0000/66/000/000/0309	ABSTRACT

(U) Several compositions of base lubricating oil with ash-free resinous additives (SB-3, DP-11, VN-18) and ash-containing additives (SK-11, DS-11, VNII NP-360) were tested in a YanZ-236 diesel engine using a 150 hour test duration. The tests involved inspection of engine parts for the extent of wearout, examination of oil composition, and piston tolerance. In general, the additive containing oils showed superior performance as compared with the pure base oil. Orig. art. has: 4 figures, 4 tables.

DATA SHEET PAGE		SUBJECT AREA	
PT5000912	20. Topic Title lubricating oil, lubricant property, lubricant additive	1. Accession no.	2. Report No.
a. INITIAL SELECTION OF OPTIMUM ADDITIVE COMBINATIONS FOR MOTOR OILS ON A BEACH WITH A SINGLE-CYLINDER AIR-COOLED ENGINE		3. Date of info	4. Date of info
		1989-07-16	1989-07-16
		5. SOURCE	6. INVENTOR
b. VSEGOVNOYE SODESHCHENIYE PO KHMILI I PRIMENENIYU PRISADOK K MASLAM I TOPIL'YU (RUSSIAN)		7. DOCUMENT NO.	8. INVENTOR NO.
2D, 1986, PRISADOK K MASLAM. TRUDY (RUSSIAN)		PTD-ET-23-149-12	FTD-ET-23-149-6
c. SECURITY AND DOWNGRADING INFORMATION		9. SECURITY CLASSIFICATION	10. COUNTRY, ORIGINATOR
		UNCL	UNCL
d. COUNTRY, NATION		11. COUNTRY	12. SECURITY CLASSIFICATION
		UR	UNCL
e. PUBLICATIONS		13. PUBLICATION DATE	14. PUBLICATION DATE
f. NEW ADD. NO.		15. TYPE PROPERTY	16. TYPE PROPERTY
P-33657-68-D-0856-P002		65-AT7021056	91-00 TRANSLATION
g. STEP NO.		17. ACCESSION NO.	18. SUBJECT
02-UR/0000/66/000/0297/0500		02-UR/0000/66/000/0309	ABSTRACT

(U) A new testing unit for lubricating oils was developed for rapid (10 hour) evaluation of oil quality, and grade. The unit based on a one-cylinder Ulyanovets Model UD-1 air-cooled engine is described in detail. The method, called IM-10FG, makes it possible to assess rapidly the composition of a given lubricating oil, to classify it according to the Premium, Heavy Duty, or Series I nomenclature, and to define optimum contents of various additives. Tables of recommended additive compositions were compiled from the test results. Oils C; these compositions were then tested on a full-scale YaMZ-407 automobile engine. Orig. art. has: 5 tables.

DATA SHEET		TEST METHOD	
CONTRACT NO.	RECEIVED BY	TESTER	TESTER
TPD00094 TWO-STROKE OIL AND OIL-ADITIVE SELECTION AND USE-PROPERTY MOTOR RATING FOR TWO-STROKE GASOLINE ENGINES	11, 21 12-AUTOMO-2 AUTOMO FILIPOV, V. P.; 16-BURULOV, V. N.; 16- 16-AUTOMO GAVRUKHIN, V. N. 13-SECURITY AREA 14-SECURITY NUMBER 15-SECURITY DATE 16-SECURITY NO. 17-SECURITY DATE 18-SECURITY AREA 19-SECURITY NUMBER 20-SECURITY DATE	1889 0539 F33657-68-D- 0866 BON2 TPP 02-UR/0000/66/000/000/0319/0324	1889 0539 F33657-68-D- 0866 BON2 TPP 02-UR/0000/66/000/000/0319/0324
UNCL. 0	UNCL. 0	UNCL.	UNCL.
TEST RECD. NO.	TEST-DESCRIPTION	TEST-DESCRIPTION	NO. OF PAGES
1889 0539	TEST ACC. NO.	TEST-NAME	10
F33657-68-D- 0866 BON2	65-ATZ021038	QU-00	TEST-NAME TEST-NAME
TPP 02-UR/0000/66/000/000/0319/0324	CONTRACT	CONTRACT	CONTRACT

(U) A mineral lubricating oil testing unit (the UTM-6-NATI) based on a one-cylinder engine and a modification of it (the UTM-6-R-NAVY) are described in detail. A reliable, short (120 hr.) testing procedure is described for rapid differentiation among the oils; heavy duty, Series 1, and Series 2. A large number of domestic and foreign oils of various quality were examined and the results of these examinations were found to agree well with the actual quality of the oils as well as with the results of the old-style 500-800 hr tests. Orig. art. has: 1 figure, 7 tables.

(U) A new versatile method (cylinder temperature 170-230 degrees C., maximum temperature of base oil 180-200 degrees C., maximum temperature of additive-containing oil 250 degrees C.) of testing lubricating oils is described. The method's reliability is assessed on the reproducibility of the solid matter within plus or minus 10 percent. The method is recommended for use in making up lubricating oils for research purposes, and for setting industrial standards for lubricating oils. Orig. art. has: 2 figures, 2 tables.



DATA SHELLING PAGE		DATA SHELLING PAGE	
61-ACCESION NO.	62-SECURITY LSC	61-ACCESION NO.	62-SECURITY LSC
TT5000017		TT50000918	
60-TITLE ADDITIVE MIXES FOR STABILIZATION OF TRANSFORMER OILS	Lubricant passivator, transformer oil, antioxidant additive	60-TITLE INVESTIGATION OF ADDITIVES AND ADDITIVE OILS BY A POTENTIOMETRIC METHOD	titrimetry, lubricant additive, oil
62-SUBJECT AREA		62-SUBJECT AREA	
11		11	
63-AUTHOR CO-AUTHORS	SHAMKOVICH, N. I.	63-AUTHOR CO-AUTHORS	LIREKHA, V. S., 16-BURDENOK, L. N.
63-AUTHOR VERSOYUZNOYE SOVESHCHANIE PO KHIMII I PRIMENENIYU PRISADOK K MASLAM I TOPLIVAM 2D, 1966, PRISADKI K MASLAM. TRUDY (RUSSIAN)	FED-HT-25-1406-68 FED-HT-25-1406-78 722501-78	63-AUTHOR VERSOYUZNOYE SOVESHCHANIE PO KHIMII I PRIMENENIYU PRISADOK K MASLAM K TOPLIVAN 2D, 1966, PRISADKI K MASLAM. TRUDY (RUSSIAN)	FED-HT-25-1406-68 FED-HT-25-1406-78 722501-78
64-SECURITY AND SOURCE/ARMED INFORMATION	UNCL	64-SECURITY AND SOURCE/ARMED INFORMATION	UNCL
UNCL	NONE	UNCL	NONE
70-REEL FRAME NO.	77-CHARTES	70-REEL FRAME NO.	77-SUPERSEDES
1889 0542	70-GRAPHICAL AREA TR	1889 0543	70-CHANGES
725657-68-D-0866 P002	X SEP ACC. NO. PUBL. SHINE RATE: 94-00	725657-68-D-0866 P002	X SEP ACC. NO. PUBL. SHINE RATE: 94-00
REV. NO.	65-ATT021048	REV. NO.	TRANSLATION NONE
REV. NO.	02-UR/0000/66/000/000/0379/0388	REV. NO.	ABSTRACT
ABSTRACT			

(U) The effectiveness of disalicylideneethylenediamine (deactivator), anthranilic acid (passivator), and phenyl-beta-naphthylamine and 2,6-di-tertiary-butyl-4-methylphenol (antioxidants) and their combinations in the stabilization of transformer oils was studied under static laboratory conditions (44 hr oxidation with oxygen in the presence of copper and iron at 100 degrees C and 49 kilovolt per centimeter potential difference). Excellent stabilization of the transformer oils was achieved by compositions containing 0.05 wt percent anthranilic acid and 0.05-0.2 wt percent antioxidant additives. Orig. art. has: 3 figures, 1 table.

(U) The applicability of the potentiometric titration method (up to 520 mV at pH equals 10) for determining the deterioration (alkalinity, metal content, ash content) of additives in oils and additive oils was investigated. It was found that all oil and oil additive deterioration indices could be determined by the potentiometric titration technique with an accuracy as high as 0.02-0.25 percent. The study revealed that imported oils with additives have higher phosphorus and zinc content than the corresponding domestic oils. In general, the potentiometric method was rated very useful and reliable. Orig. art. has: 7 figures, 5 tables.

K. I. Ivancov, Ye.L. Vilyanskiy, A.A. Zinetskiy and K.K. Akhmedarov, Use of Paired Anticorrosion Oils . . . . .	45
M.I. Shakinovich, Additive Mixes for Stabilization of Transformer Oils . . . . .	1
V.S. Luneva and L.M. Burdenuk, Investigation of Additives and Additive Oils by a Potentiometric Method . . . . .	16
<b>TABLE OF CONTENTS</b>	
S.E. Kravyn, Yu.M. Shechter, N.K. Levitin, A.V. Vilenskin, V.F. Kalashnikov, A.L. Dvortserg, V.I. Goryacheva, L.G. Khoroshilova, and R.I. Bessmerity, Corrosion-Inhibitor Additives and Preservative Running Oils Based on Them . . . . .	1
O.S. Oblyukhova, V.V. Protasov, T.T. Kiseleva, and G.V. Burtseva, Operating Experience with ZIL-130 Engines Using AS-8011 with VNII NP-36G Additive . . . . .	16
N.I. Sigal, Ye.P. Slobov, V.N. Yefremov, G.M. Gusev, G.P. Belianchikov, I.A. Trakhtonenko, I.A. Kozlomov and N.V. Versnov, Test Results for Oils with Additives Used in the YaIZZ-238NE Engine . . . . .	23
C.S. Oblyukhova and N.H. Kryucheshnikova, Effectiveness of Using Additives in Hydraulic-Coupling Oils for Passenger Automobiles . . . . .	30
Ye.G. Semenido, I.A. Trakt'yenko, N.V. Shatogolev, N.A. Senichkin, A.S. Lozar', and V.I. Strapov, Coordination of Ash-Forming and Polymeric Ash-Free Additives in Motor Oils as a Way to Further Quality Improvement. .	40
V.P. Filippov and S.M. Gubarev, Selection of Optimum Additive Combinations for Motor Oils on a Bench with a Single-Cylinder Air-Cooled Engine . . . . .	47
S.G. Arutyun, G.P. Belyanchikov, I.N. Parfenov, T. Khanlarov, I.R. Khodzhanov, G.K. Cheshmina, Ye.V. Firdanova, Tests of Soviet and Foreign Motor Oils in the TIM-6-MAT and CIM-6M-MAT Single-Cylinder Bench Installations . . . . .	62
V.P. Filippov, V.M. Bokolov and V.V. Davydov, Oil and Oil-Additive Selection and Proper Motor Rating for Two-Stroke Gasoline Engines . . . . .	76
A.I. Fazov and E.S. Zuskev, In-Storage Property Changes of Oils of Various Grades and Determination of Their Purity. . . . .	85

inhibitors that effective inhibiting additives can be obtained from nitrated oils that have been subjected to special treatment. In developing a technology for production of these additives, it was first necessary to solve such problems as selection of the raw material for nitration and the nitrating agent, as well as finding the optimum conditions for nitration and final treatment of the product. Two-stage reactions take place when mineral oils are nitrated:

nitration proper

$\text{RAr}-\text{HNO}_2, \text{H}_2\text{SO}_4,$

oxidation

$\text{RAr}-\text{HNO}_2, \text{KOH/H}_2\text{O}$

where RAr are the alkylaromatic and naphthalenoaromatic hydrocarbons of the oil.

#### CORROSION-INHIBITOR ADDITIVES AND PRESERVATIVE RUNNING OILS BASED ON THEM

S.E. Kreyv, Yu.N. Shekhter, M.M. Levitin, A.V. Viletskin,  
V.P. Kalesnikov, A.L. Solberg, V.I. Goryacheva,  
L.D. Khorozhilova, and K.I. Bessmertny

Tractor-operating experience has shown that engine and transmission parts are subject to rusting during operation and storage under the influence of moisture and acid products that have entered the lubricating oil or been formed in it. Since the commercially available motor and transmission oils have unsatisfactory protective properties [1], and if used as preservative oils of types F-17, NO-203 etc., there is a need for development of preservative running oils. An effective protective and preservative running oil is an aromatic nitrated protective additive. Special additives that act as corrosion inhibitor must be added to the existing commercial oils. In addition to the additives that improve operational properties, use of preservative running-inhibiting oils will shorten the service lives of motor-traction equipment and ultimately save a substantial amount of money.

The recommended protective additives are oil-soluble corrosion inhibitors: the reaction product of thiophosphoric acids with quinoid compounds and other thiophosphorus-containing substances [2], various derivatives of amine [3], organic acids, their esters and salts [4], derivatives of sulfonic acids, nitrated products [7], etc.

It has been established by laboratory studies of the protective and use properties of oil with various corrosion

Table 1. Physicochemical Properties of Nitrated Oils  
(Nitration with 60% Nitric Acid at 15-55°C for 4 h)

a Number	Frances 3	AS-4.5	AS-3	E-4	E-11	Sto
Frances type, deg C (nitration temp.)	10	15	15	15	15	15
Frances, %, min	63	65	62	62	62	65
Frances in naphthalene, PPN, % KOH/E	...	...	...	...	...	...
Frances in benzene, PPN, % KOH/E	...	...	...	...	...	...
Frances viscosity, cSt (temperature 100°C, oil temperature 20°C)	95	62	36	10.1	7.4	5.9
Frances viscosity, cSt (temperature 20°C, oil temperature 20°C)	90	82	17.6	17.3	15.0	6.67
Frances viscosity at 100°C, cSt	-	-	6.75	6.93	-	-

KEY: (a) indicators; (b) nitrated oils; (c) spiral; (d)  
(1) AS-4.5 (e) Frances; (f) E-4; (g) Stoen; (h) Sto;  
(i) PPN formed during nitration; (j) oil; (k) none;  
(l) ash; (m) Indicator alkalinity, mg of KOH/E; (n)  
phenolphthalein; (o) neutralization of bromphenol blue;  
(p) viscosity at 100°C, cSt; (q) nitrogen content, %.

Various oxygen-containing compounds are obtained in a nitrating oil. The oxidation reaction: as is, in thiacids, lactones, esters, etc., and they are capable of the following secondary reactions:  
1) condensation (polymerization), extending to the formation of asphalt-like substances;

b) nitration, since the oxygen-containing compounds of the oil nitrate considerably more readily than the initial oils (the latter are subject to nitration or previous oxidized oils).

The brown nitrogen-oxide  $\text{NO}_2$ ,  $\text{N}_2\text{O}_4$ ,  $\text{NO}$ ) vapors evolved during the oxidation reaction in turn promote oxidation of the hydrocarbons. Thus the oxidation process of the oil may become autocatalytic if substantial amounts of nitrogen-oxide vapors are liberated.

[\*] Is known that polycyclic aromatic hydrocarbons with short side chains and tarry substances are oxidized preferentially in mineral oils, undergoing secondary condensation reactions with formation of asphalt-like products. For nitration, therefore, we selected oils from which these compounds had been removed by selective refining. Oils AS-9.5, DS-8 and DS-11 obtained from eastern petroleums gave the best results (Table 1).

The direction of the nitration process and its degree of conversion can be regulated by applying various nitration methods to the mineral oils.

#### The following oil-nitration processes were examined.

1. Nitration with nitric acid at various concentrations (from 10 to 98%), with various acid-to-oil proportions. The optimum nitrating formulas were found in each case: addition and standing times for the acid, nitration temperature, etc. As the nitric acid concentration increases, the content of nitro compounds in the nitrated oil increases without any marked increase in the content of oxidized products. Nitration with weak acids intensifies the oxidation reaction.

2. Nitration with nitrating mixtures or sodium nitrate or nitrite in the presence of sulfuric acid. These methods enable us to nitrate oils with formation of mononitro compounds, minimizing the oil-oxidation reaction.

3. Nitration with catalysts. By the use of various catalysts, the process can be controlled either in the direction of nitration (sodium nitrite, zinc stearate) or in the direction of oxidation (mercury salts).

First, the combined nitration and oxidation products were extracted from the nitrated oils and then separated by a secondary gasoline extraction.

It was shown that the nitro compounds are those principally responsible for the high protective, detergent and other properties of the nitrated oils. The amounts of nitrogen in them, determined by a specially developed method (a modified Kielbahl method) [8], indicate that mononitro compounds (nitrogen content 3.7-3.9%) are formed during nitration under normal conditions.

The nitro group is found attached to the benzol ring (or, more rarely, to the naphthenic nucleus), for the most part at the para-position with respect to the alkyl radical.

Although they are less effective than the nitrocompounds, oxygen-containing products accompanying them in certain proportions - approximately 0.5:1 - form synergistic mixtures that exhibit the highest protective properties. In nitration, therefore, it is not desirable to produce only nitro compounds, for example, by nitration with anhydrous nitrating mixtures.

AS-9.5 oil nitrated with 40% nitric acid at 60°, cone. nitration was chosen as the base for the oil and fuel corrosion-inhibiting additives. This oil contains 6-7% of nitro compounds and 4-5% of oxidation products.

After carrying out nitration and eliminating the spent nitric acid, the acidic nitrated oil (acid number 20, mg of KOH/g) must be rendered slightly alkaline with some sort of neutralizing agent. Salts of sodium, calcium, and magnesium, and especially those of lead and aluminum, showed good protective, detergent, and dispersing properties [9].

Alkalinized nitrated oils show high ash contents and excessive alkalinity, which indicates not only the presence of salts of the type  $(\text{HArCOO})_2\text{Ca}$  in the nitrated oil, but also that they contain a finely dispersed colloidal suspension of  $\text{CaO-Ca}(\text{OH})_2$  and perhaps also  $\text{CaCO}_3$ , stabilized by surface-active compounds.

Work was done to study the mechanism of action of the nitrated oils and the products based on them (NG-204 and Ni-204 liquid lubricants and ATOR-1 corrosion-inhibiting additive). It was established that nitrated oils and products based on them sharply reduce the permeability of the oil films to moisture and vapor [9] and reduce (at a certain concentration, to zero) the surface tension at the oil-water and water-(adsorbed oil film) interfaces.

They increase the wetting angle of an oil drop on water and reduce the wetting angle of a water drop on the oil's surface. This is particularly conspicuous with the ANOR-1 additive, and for lead and aluminum salts of the nitrated oil, i.e., in the most effective corrosion inhibitors. The ability to displace water from the surface of the metal and block its passage through the hydrophobic adsorption films that form is of great practical importance, since it permits the use of nitrated oils to preserve wet surfaces, as well as in varnishes, primers, paints, watered fuels, etc.

Studies conducted in a special chamber by using electric current for external polarization of plates [1] indicated that sodium and calcium salts of nitrated oils showed distinctly the properties of oil-soluble anodic atmospheric-corrosion inhibitors, while the lead and aluminum salts are adsorbed on both the anodic and cathodic areas of the metal exposed to corrosion, and

the protection given these areas is of the screening type.

Two problems were faced in developing corrosion-inhibitor additives for the motor and transmission oils.

#### 1. Development of a corrosion-inhibitor additive which,

in addition to the base oils ~ oils containing a combination of additives, will endow these oils with high protective properties without any deterioration of their motor properties.

#### 2. Creation of a multifunctional corrosion-inhibiting additive that would, on addition to the base oils, endow them with high protective properties and simultaneously improve their wetting and anticorrosion properties.

The nitrated oils themselves or their salts (sodium, calcium, or aluminum) may be used as additives of the first type. Nitrated oil will also be taken as a base for creation of multifunctional additives of the second type. However, acidic nitrated oils are alkalized in the presence of special promoters to improve the detergent and dispersing properties of the additives.

Like the use of special technological devices (single fast evaporation of water, passage of carbon dioxide, etc.), the presence of promoters during alkalization results in a sharp increase in the ash figure and alkalinity of the oil solution of the additive and, as a consequence, improvement of its wetting properties.

Promoters that have been tested and are recommended by the authors include alkylphenols, arylamines, alcohols, synthetic fatty acids, hydroxyethylated products, and many other compounds.

After appropriate refinement, therefore, the following technology was adopted for product of AKR-1 additive: selectively refined DS-6 or RS-11 oils are treated with 30% of 10% nitric acid, followed by addition of 10% stearic acid; the mixture is neutralized with 20% calcium oxide, heated to 120°C to eliminate water, and centrifuged. Two industrial batches of the additive were prepared by the above technology at a pilot plant at the "Neftegaz" refinery. The material balance of this process is given below.

Taken, % Yield, %

Production of nitrated oil	
Mineral oil DS-8.....	169
Nitrated oil.....	103
Volatile products.....	3
Spent nitric acid (30%) .....	26
Mechanical impurities.....	4
Litter.....	3
TOTAL.....	133
TOTAL.....	130

Production of AKR-1 Additive	
Nitrated oil.....	100
AKR-1 Additive.....	112
Waste.....	28
Calcium oxide.....	10
TOTAL.....	130

Several laboratory batches of the AKR-1 additive were prepared substituting synthetic fatty acids for the stearic acid. It was established that the synthetic fatty acids combine well with the nitrated oil (Table 2) and that the resulting additives are highly effective (Table 3).

The configuration of the apparatus and the flow chart for AKR-1 production at the "Neftegaz" refinery are examined below.

Mineral oil (Fig. 1) and nitric acid are fed continuously into reactor 1. Heat is liberated as a result of the nitration reaction and removed by cold water. The reaction mixture enters reactor 2, where the nitration reaction continues for another h at 55°C. Then the mixture is directed into settling tanks 3 and 4. The spent acid drains by gravity into apparatus 5 and is reused for nitration. The acid nitrated oil goes into apparatus 6, where it is mixed with melted stearin (or synthetic fatty acids [SPA] (GK)) and neutralized with calcium oxide (GK). The product is then dehydrated in the system consisting of heat exchanger 9 and column 10. The product flows from the bottom of column 10 into centrifuge 12, is heated in tank 11, and further purified on the ultracentrifuge 15, from which the final additive is run into the shipping containers.

Table 2. Physicochemical Properties of AKR-1 Additive Obtained using Synthetic Fatty Acids

Additive	Properties of AKR-1 Additive			
	Indicator of alkalinity of base oil	Specific gravity at 20°C	Viscosity at 100°C, mm <sup>2</sup> /s	Indicative alkalinity of base oil
Stearic acid	55	0.90	35	65
Calcium stearate	55	0.93	27.9	75
Stearic acid + SPA	53	0.93	26.4	64
Stearic acid + SPA + calcium oxide	53	0.93	26.4	64
Stearic acid + SPA + calcium oxide + calcium carbonate	53	0.93	26.4	64

Key: (a) Indicator; (b) synthetic fatty acids used; (c) broad fraction from Mendeleyev plant; (d) C<sub>18</sub>-C<sub>1</sub> fraction from Lebedev plant; (e) C<sub>18</sub>-C<sub>1</sub> fraction from Volgograd refinery; (f) C<sub>18</sub>-C<sub>1</sub> fraction; (g) ash; (h) total; (i) sulfate; (j) indicator alkalinity, m; of FOH(G); (k) phenolphthalein; (l) bromophenol blue; (m) viscosity at 100°C, etc.



Concentrations of various organic acids in 10% KOH/a					
Concen-	Concen-	Concen-	Concen-	Concen-	Concen-
tration %	concentration in 10% KOH/a	concen-	concen-	concen-	concen-
3.8	2.3	46.7	26.1	14 cytos (G-3 octa-	r
0.9	-2.9	6.9	19.9	26 cytos (G-3	r
3.4	6.6	55.0	30.3	50 cytos (G-3 octa-	r
1.8	2.5	16.3	53.0	90 cytos (G-3 octa-	r
1.0	-5.3	18.4	14.4	Cytos (G-3 octa)	r
3.0	-1.2	46.9	31.3	60 cytos (G-3 octa)	r
4.0	0.9	49.9	43.4	14 cytos (G-3 octa)	r
1.0	-5.2	5.3	25.0	Cytos (G-3 octa)	r
3.75	2.5	52.7	58.5	78 cytos (G-3 octa)	r
1.7	4.4	11.7	35.0	23 cytos (G-3 octa)	r
4.5	4.7	54.3	122.3	46 cytos (G-3 octa)	r
3.7	2.5	44.8	53.4	78 cytos (G-3 octa)	r
				(G-3 octa) 2	

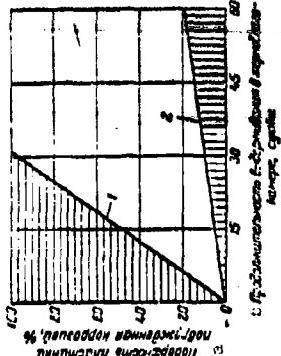


Fig. 2. Curves of time increase of corrosion damage to steel plates surface (G-4 heat and humidity chamber, 7 h at 50°C, 17 h at 20°C; humidity 100%, plate material St. 3). 1) Commercial oil; 2) commercial oils + 10% AKOR-1.

KEY: (a) area of plate damaged by corrosion, %; (b) holding time in heat-and-humidity chamber, days.

an explosion-proof apparatus and an additional installation for driving off the gasoline. However, the cost of building this apparatus is fully covered by the high quality of the resulting product, which is superior to a number of indicators to foreign additives used for similar purposes.

The protective properties of motor and transmission oils containing AKOR-1 additive in various concentrations were studied under laboratory conditions. For this purpose, metal plates - steel, cast iron, bronze, brass, copper, lead and other non-ferrous metals - were coated with thin films of oil containing the additive and placed in water. In humidity chambers, or in chambers containing corrosive environments. In all cases, corrosion appeared on the plates much later than when the plates were protected with oils not containing additives [1, 10]. By way of example, Fig. 3 shows generalized time curves of the corrosion damage to steel plates in a heat-and-humidity chamber. When they were protected with commercial oils, the damage curve rises sharply, and plates coated with oils having better protective properties are completely corroded in no more than 30 days. The difference in the protective properties of the oils before and after addition of the various functional additives is illustrated by the width of the shaded region in Fig. 2. The increase in the area of the corrosion centers after addition of 10% of AKOR-1 additive to the oil is much slower, and no more than 20% of the plate area has been corroded after storage for two months in the chamber.

- (k) viscosity at 100°C, cst
- (u) SFA, ...fracture
- (l) solubility in oils
- (v) ...days (small amount of deposit)
- (m) 10% in MT-16
- (n) same
- (o) 10% in transformer oil
- (p) one day (deposit)
- (q) AS-9.5
- (r) DS-11
- (s) stearic acid
- (t) calcium oxide hydrate
- (u) ...days (without deposit)
- (v) sodium oxide hydrate
- (w) one day (deposit)

Table 4. Evaluation of Protective Properties of Oils in Exhaust-Gas Chamber (30 min at 50°C, 2 h at 20°C)

a No.	b Oxidative stability, min	c Ash content, %	d Induced ash content, %	e Corrosion, %	f Viscosity, cSt	g Viscosity, cSt
b AKO-11	150	0	0	—	—	—
1 MT-11 + 10% AKOP-1	50	0	0	—	—	—
MT-16	70	0	0	—	—	—
q MT-16 + 10% AKOP-1	50	0	0	—	—	—
r MT-16	90	0	0	—	—	—
t MT-16 + 10% AKOP-1	10	0	0	—	—	—

KEY: (a) oil; (b) condition of nonferrous-metal plates after testing; (c) bronze; (d) surface condition; (e) amount of surface affected by corrosion; (f) brass; (g) aluminum; (h) DSP-11; (i) brown deposit; (j) reddish-brown point foci; (k) moderate darkening of entire surface; (l) DSP-11 + 10% AKOR-1; (m) no changes; (n) point foci barely discernible; (o) dark brown point foci; (p) dark point foci; (q) MT-16 + 10% AKOR-1; (s) minute dark point foci; (t) MT-16 + 10% AKOR-1.

Table 4 presents certain data characterizing the protective properties of oils with the AKOR-1 additive for nonferrous metals.

The high protective properties of oils containing AKOR-1 additive have been confirmed by the results of storing automotive engines and transmissions on an outdoor platform (Table 5).

Certain physicochemical and operational properties of the oils undergo changes after introduction of AKOR-1 (Table 6). Thus, the viscosity level of the oil at 100°C rises slightly and ash content increases. The detergent and anticorrosion properties of the oil are improved substantially.

The protective additive has no noticeable detrimental effect on the quality indicators of the motor oils and transmission lubes.

Table 5. Surface Condition of Parts, Subassemblies and Units Prepared for Preservation With the Use of Various Commercial Oils with AKOR-1 Additive

a No.	b Preserved surfaces, units and assemblies	c Storage time, months	d Condition of surfaces	e Corrosion, %	f Type designation	g Storage time, months
e Automobile type MT-16	—	—	—	—	—	—
f Gasoline, kerosene, mineral oil, water	—	—	—	—	—	—
g Diesel engine	—	—	—	—	—	—
h Gasoline engine	—	—	—	—	—	—
i Diesel engine	—	—	—	—	—	—
j Gasoline engine	—	—	—	—	—	—
k Gasoline engine	—	—	—	—	—	—
l Gasoline engine	—	—	—	—	—	—
m Gasoline engine	—	—	—	—	—	—
n Gasoline engine	—	—	—	—	—	—
o Gasoline engine	—	—	—	—	—	—
p Gasoline engine	—	—	—	—	—	—
q Gasoline engine	—	—	—	—	—	—
r Gasoline engine	—	—	—	—	—	—
s Automobile type MT-16	—	—	—	—	—	—
t Gasoline engine	—	—	—	—	—	—
u Gasoline engine	—	—	—	—	—	—
v Gasoline engine	—	—	—	—	—	—
w Gasoline engine	—	—	—	—	—	—
x Gasoline engine	—	—	—	—	—	—
y Gasoline engine	—	—	—	—	—	—
z Gasoline engine	—	—	—	—	—	—
aa Gasoline engine	—	—	—	—	—	—
bb Gasoline engine	—	—	—	—	—	—
cc Gasoline engine	—	—	—	—	—	—
dd Gasoline engine	—	—	—	—	—	—
ee Gasoline engine	—	—	—	—	—	—
ff Gasoline engine	—	—	—	—	—	—
gg Gasoline engine	—	—	—	—	—	—
hh Gasoline engine	—	—	—	—	—	—
ii Gasoline engine	—	—	—	—	—	—
jj Gasoline engine	—	—	—	—	—	—
kk Gasoline engine	—	—	—	—	—	—
ll Gasoline engine	—	—	—	—	—	—
mm Gasoline engine	—	—	—	—	—	—
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tt Gasoline engine	—	—	—	—	—	—
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oo Gasoline engine	—	—	—	—	—	—
pp Gasoline engine	—	—	—	—	—	—
qq Gasoline engine						

[Key to Table 5, cont'd.] compressor, valves; (g) 90% DS-8 + 10% AKOR-1; (h) 21 months (no corrosion); (i) thin unbroken oil film; (j) transmission; (k) surfaces of gear teeth; and shaft splines; (l) 90% MR-16p + 10% AKOR-1; (m) same; (n) YaAZ-206-A engine; (o) highly finished surfaces of engine cylinders, timing gear, compressor; (p) 19 months (no corrosion); (q) YaAZ transmission; (r) surfaces of gear teeth, shafts, guide rollers, shifting cylinder and detents; (s) GAZ-51 engine; (t) honed cylinder surfaces; (u) 90% ASp-0.5 + 10% AKOR-1; (v) GAZ-51 transmission; (w) 90% nigrol + 10% AKOR-1; (x) GAZ-59 engine cylinder blocks; (y) honed surfaces, parting planes; (z) 70% DS-8 + 10% AKOR-1; (aa) 20 months (no corrosion); (bb) 90% AS-9.5 + 10% AKOR-1; (cc) unbroken oil film, streaks of oil on honed surfaces; (dd) 22 months (no corrosion); (ee) unbroken oil film; (ff) GAZ-69 transmission; (gg) 90% TAP-15 + 10% AKOR-1.

Table 6. Physicochemical and Operational Properties of DSp-11 Oil with 10% of AKOR-1 Additive

Parameter, com pounds and elements	Concen-									
	a	b	c	d	e	f	g	h	i	j
1. Temperature, °C	105	115	105	115	115	115	115	115	115	115
2. Viscosity, cSt	0.05	0.05	0.05	0.05	0.05	0.05	0.05	0.05	0.05	0.05
3. Density, g/cm <sup>3</sup>	0.86	0.86	0.86	0.86	0.86	0.86	0.86	0.86	0.86	0.86
4. Viscosity index	125	125	125	125	125	125	125	125	125	125
5. Temperature, °C	22	22	22	22	22	22	22	22	22	22
6. Temperature, °C	35	35	35	35	35	35	35	35	35	35
7. Maximum operating temperature, °C	55	55	55	55	55	55	55	55	55	55
8. Maximum operating temperature, °C	113	113	113	113	113	113	113	113	113	113
9. Flash point, °C	40	40	40	40	40	40	40	40	40	40
10. Pour point, °C	-45	-45	-45	-45	-45	-45	-45	-45	-45	-45
11. Thermal-oxidation stability, min;	160	160	160	160	160	160	160	160	160	160
12. Potential detergency, %	—	—	—	—	—	—	—	—	—	—

KEY: (a) indicator; (b) oil without additive; (c) additive; (d) nitrated DS-8 + 10% stearin + 10% Ca(OH)<sub>2</sub>; (e) nitrated AS-9.5 + 10% stearin + 5% Ca(OH)<sub>2</sub>; (f) nitrated AS-9.5 + 10% SPA (C<sub>10</sub>-C<sub>10</sub>) + 10% Ca(OH)<sub>2</sub>; (g) nitrated AS-9.5 + 10% SPA (C<sub>10</sub>-C<sub>10</sub>) + 10% NaOH; (h) nitrated AS-9.5 + 10% SPA (C<sub>17</sub>-C<sub>20</sub>) + 5% Ca(OH)<sub>2</sub> + 5% NaOH; (i) nitrated AS-9.5 + 10% SPA (C<sub>1</sub>-C<sub>10</sub>) + 3% Ca(OH)<sub>2</sub> + 2% NaOH; (j) viscosity, cst; (k) at; (l) ash; (m) temperatures, °C; (n) flash point; (o) pour point; (p) thermal-oxidation stability T<sub>200</sub>, min; (q) potential detergency, %; [Key cont'd. on page 14]

[Key to Table 6, cont'd.] (r) Detergent power on PZV machine, points; (s) carbon scaling at 400°C; (t) anticorrosion properties on PZV machine at 200°C (lead).

The running properties of motor oils with the AKOR-1 additive were checked during bench tests on 0D-9 (a single-cylinder forced diesel), GAZ-69, YaAZ-238, 2Ch-8, 5/11 and Ural-375 engines and use tests in GAZ-51, ZIL-157 and KRAZ-214 motor vehicles. It was reported in all cases that the engine parts remained cleaner and showed practically unchanged wear after addition of AKOR-1.

### CONCLUSIONS

1. AKOR-1 protective additive was prepared from nitrated oils that had been treated with sodium or calcium oxide hydrates or mixtures thereof in the presence of stearic acid, SPA, and other compounds.

2. A flow chart for production of the additive was worked out and used.

3. AKOR-1 additive effectively improves the ability of commercial motor oils and transmission lubes to protect steel, cast-iron and nonferrous-metal parts from rusting.

4. The detergent and anticorrosion properties of oils with AKOR-1 additive are superior to those of oils without this additive, and other properties show virtually no differences.

5. In pure form or diluted with light oil, AKOR-1 additive can be used on a par with liquid preservative oils for internal preservation of mechanisms.

6. AKOR-1 additive, added to motor oils and transmission lubes in a concentration of 10%, has successfully passed use tests:

a) for protective properties when used to preserve engines and motor-vehicle transmission assemblies during storage or an open-air platform for more than 20 months;

b) for operational properties in bench tests on 0D-9, 2Ch-8, 5/11, GAZ-69 and YaAZ-238 engines and use tests in GAZ-51, ZIL-157 and KRAZ-214 motor vehicles.

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#### OPERATING EXPERIENCE WITH ZIL-130 ENGINES USING AS-8 OIL WITH VNIIP MP-360 ADDITIVE

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T.T. Kisileva, and G.V. Burtseva

DS-8 oil with VNIIP MP-360 additive [VNIIP: All-Union Scientific Research Institute of Oil and Gas Refining and Production of Synthetic Liquid Fuels], which had been recommended after long-term (600-hour) bench tests for V-type curbed truck engines, was taken for extended road-use tests. Since 1964, this oil has been produced according to GOST 10541-63 under the type designation AS-8.

The tests were conducted with 70 vehicles in various climatic zones of the country (Crimea, Stavropol', Orel, Pskov) on a year-round basis, on hard-paved mountain and flat lands highways. Vehicles of the Crimea fleet were tested under routine operating conditions on the Yalta-Simferopol' and Yalta-Sevastopol' routes. The vehicles of the other fleets were run under routine operating conditions for 10-30% of the total mileage, and used for the remaining time in tire tests. The conditions of use are given in Table 1. A-76 gasoline was used in all fleets during the tests.

Plans called for replacing piston rings during the tests when oil burnoff had reached 0.8 liter/100 km and the bearing inserts when the oil pressure in the system had dropped to 1.9 kg/cm<sup>2</sup> at 1000 rev/min.

At the end of the tests, when the vehicles had accumulated 108-213 thousand km, these oil-consumption and pressure limits

Table 1. Vehicle Operating Conditions

oil-change Intervals.

## TEST RESULTS

a	Astrodon № 1122. F. Crassipes . . . .	C. Experiments done in laboratory		C. Experiments done in field		C. Experiments done in field		C. Experiments done in field	
		1	2	3	4	5	6	7	8
		Hatching, m.	80	40	40	55—65 (in normal tempera- ture)	50	700— 750	700— 750
		Incubation, m.	30	—	—	40—55 (in normal tempera- ture)	—	—	—
		Development, m.	22	350	—	—	—	—	—
		Incubation, m.	22	350	—	—	—	—	—
		Hatching, m.	21	350	—	73	22	40—55 (in normal tempera- ture)	40—55 (in normal tempera- ture)
		Incubation, m.	21	350	—	—	—	—	—
		Development, m.	21	350	—	—	—	—	—
		Incubation, m.	21	350	—	—	—	—	—
		Hatching, m.	21	350	—	—	—	—	—

**Note:** Servicing intervals: TO-1 every 1600–2000 km; TO-2 every 6000–8000 km [TO: technical maintenance].  
**TO-1:** (a) fleet; (b) number of vehicles; (c) road characteristics; (d) length, km; (e) concrete, %; (f) asphalt, %; (g) traprock and gravel, %; (h) load, tons; (i) speed, km/h; (j) technical average; (k) topographic; (l) average mileage, km; (m) convoy No. 11202, Stavropol; (n) 55–65 (on asphalt), 40–55 (on mountain and traprock roads); (o) convoy No. 1120, Omsk; (p) 51 (on asphalt), 40 (on gravel); (q) conveyor number 114, Orel; (r) 57–65 (on asphalt).

had not been reached. However, the bearing inserts of almost all vehicles were replaced after 100-150 thousand km in order to prolong crankshaft life.

Piston rings were replaced at mileage of less than 150 km on only 12 engines of the Stavropol fleet, in to ensure full performance for the tire tests.

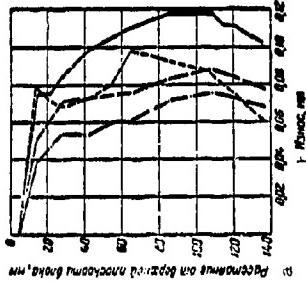
The following work was done at the end of the tests to rate the use properties of the oil:

a) visual inspection of engine parts to determine the detergent, antioxidant, anticorrosion and low-temperature prop-

b) micrometer measurements on engine parts to determine a

c) analysis of the used oils, to permit determination of quality changes during use and adjustment of the recommended

**Condition of engines.** It was established on visual inspection to determine engine condition that AS-8 oil with NII NP-360 additive has satisfactory detergent properties. Varnish on the cooled areas of the pistons came to 5-6 points on the PTV scale. The rest of the piston-skirt surface was rated with varnish ranging in color from yellow to dark reddish brown. No burning of the rings was observed on any of the engines inspected, and the combustion-chamber carbon deposits were insignificant. The anticorrosion properties of this oil were evaluated from the surface condition of the bearing-insert anti-friction layer. There were no traces of corrosion damage on any of the inserts inspected, which indicates that the oil has satisfactory anticorrosion properties.



The small amounts of gum on the engine parts and the insignificant amounts of low-temperature deposits (sludge) that were formed indicate that AS-B oil has fully satisfied the requirements necessary for temperature resistance.

A highly important advantage of the VNIIT MP-360 additive established during the tests is that it shows little tendency to form barium-salt deposits on the engine's exhaust valves.

**Engine wear.** The antiwear properties of AS-8 oil with 'NII NP-360 additive were determined by micrometer measurements on parts 24 engines after 5,213 thousand kilometers.

Cylinder-sleeve wear after accumulation of up to 200 thousand km did not exceed 0.1 mm for most of the engines; here the average or considerably longer than in the chromed unplated upper compression sleeves wear at 0.5-1.0/1000 km.

Cylinder sleeves of the ZIL-130 engine wear over their entire height. It can be explained by the favorable circumstances without stopping, high speed, Ni-Resist insert in the top of the wear properties of the oil.

The effectiveness of using engine oils with high-quality multifunctional additives has also been affirmed by foreign investigators.

Thus, K. English [1] reports that when an engine was run on HD oil (which corresponds to AS-8 oil) with the VIII NP-360 additive at a coolant temperature of 80°C, sleeve and piston-ring wear is reduced by a factor of approximately 5 by comparison with the wear observed when the engine is run on plain oil (without additives).

Piston-ring-wear data for 11 engines are given below.

Engine mileage, thousands of km	Wear mm (over total mileage)	Wear mm (per 10 thousand km)
57.89	0.0029-0.020	0.5-2.5
108-147	0.020-0.128	1.3-9.5
150-213	0.021-0.100	1.4-4.7

Crankshaft wear was determined on the connecting-rod journals for 24 engines and the main-bearing journals for 12 engines. The connecting-rod journals were measured with the engines partly and completely dismantled, and the main bearings only after full disassembly. The results of the micrometer measurements are given in Table 2.

Table 2. Main- and Connecting-Rod Journal Wear in ZIL-130 Engine

Mileage, km	a C maximum journal wear	b Bearing wear, mm	h maximum journal wear			
			d minimum journal wear	e maximum bearing wear	f average bearing wear	g maximum bearing wear
10,000	0.010	—	0.000	0.007	0.007	0.007
10,000-20,000	0.010	0.005	0.000	0.002	0.002	0.002

KEY: (a) mileage, km; (b) journal wear, mm; (c) main bearing; (d) minimum; (e) maximum; (f) average; (h) connecting-rod bearing; (1) less than.

The wear of the connecting-rod bearing inserts at a vehicle mileage of 135 thousand km averaged 0.0065 mm, and the main-bearing insert wear 0.008 mm.

Over the indicated vehicle mileage, the average clearance between the crankshaft connecting-rod journal and the inserts with the connecting-rod assembled came to 0.073 mm, and 0.075 mm for the main bearings (the acceptable clearances for new engines are 0.026-0.065 mm).

The wear of other engine parts (connecting rod upper end, camshaft, valves, rockers, lifters, etc.) was insignificant, and it was unnecessary to replace these parts before 2(0 thousand km).

**Oil consumption.** As we know, oil consumption depends on a number of factors. K. Dzhordzhi [2] notes the following basic causes of increased oil consumption:

- a) mechanical trouble - wear of piston-ring grooves and damage to the piston skirt; ring wear, loss of ring resiliency, seating of rings in grooves too tight or too loose; burning of the compression rings or plugging of slots in oil-control rings; wear, scoring or scratching of the honed cylinder surface, etc.;
- b) operating conditions - high oil level, dilution of oil with fuel due to carburetor maladjustment or low temperature in the engine's cooling system; driving speed;
- c) oil quality - volatility, viscosity index, etc.

The above indicates how difficult it is to establish operating oil-consumption norms.

In these tests, oil consumption was recorded on the basis of the amounts drained at changes and the amounts added to replace burnoff during use.

The oil-burning rate changed insignificantly as a function of engine mileage. Thus, the average oil-burning rate was 0.3 liter/100 km up to 50 thousand km and 0.45 liter/100 km to 100 thousand km. For mileages up to 150 thousand km, oil consumption came to 0.35 liter/100 km, which is probably to be explained by piston-ring changes on some of the engines. The average consumption rate rose again to 0.4 liter/100 km at the vehicles accumulated 200 thousand km.

Control trips were taken periodically in all three to verify and adjust the specific oil-burnoff figures. Figure 2 shows average oil-burnoff values from control trips with nine vehicles that had accumulated 150-180 thousand kilometers. It follows from these data that as vehicle speed is increased from 45 to 75 km/h, the oil burnoff rate increases by a factor of 2, and by a factor of 4 when the speed is increased from 45 to 95 km/h.

The average oil-consumption figures indicated above are confirmed quite accurately by the data obtained in the control trips.

Table 3 presents over-all operating fuel- and oil-consumption figures. On the basis of these results, taking into consideration that normal vehicle operating conditions are different (short-haul runs, city traffic, extended or full-time trailer hauling, less highly qualified servicing), and the

consequent higher consumption, we would recommend lowering the oil-consumption norms for the ZIL-130 engine to 2.5% of the amount of gasoline from 3.5%, which is the value presently in force for carburetor engines.

**Change in quality of oil and the oil-change interval.** To establish rational oil-change intervals, oil specimens were taken for analysis during the tests. The analytical results showed that when the oil is changed at less than 8000 km, its 50°C viscosity has increased to 47 cst (vs. 42 for fresh oil). In some cases, fuel leakage into the crankcase during winter lowers the viscosity to 35 cst.

While the fresh oil contains no organic acids, its acid number rises to 0.44 mg of KOH/g.

The total mechanical-impurity content rises to 0.65%. This comparatively low figure indicates effectiveness of oil cleaning.

Table 3. Over-All Operational Fuel- and Oil-Consumption Figures

a Autobuses	b Weight of vehicle tonnes	Percent actual fuel consump- tion over norm		Percent actual oil consump- tion over norm		Percent actual oil consump- tion over norm	
		c Average speed km/h	d Oil addi- tions liters/100 km	e Oil addi- tions liters/100 km	f Oil addi- tions liters/100 km	g Oil addi- tions liters/100 km	
i. Autobuses № 1202. r. Gorky.	65.2	0.13	0.44	0.32	0.37	0.45	1.4
j. Autobuses № 1224. r. Krasnodar.	65.2	0.13	0.48	0.54	0.61	0.67	1.65
k. Autobuses № 1244. r. Krasnodar.	36.8	0.13	0.48	0.54	0.61	0.67	1.82
l. Autobuses № 1245. r. Orenburg.	35.6	0.13	0.39	0.45	0.52	0.58	1.47
m. Autobuses № 1246. r. Krasnodar.	65.6	0.13	0.26	0.65	0.69	0.73	1.52
n. Autobuses № 1247. r. Krasnodar.	65.6	0.13	0.26	0.65	0.69	0.73	1.72

KEY: (a) fleet; (b) actual average fuel consumption, liters/100 km; (c) oil consumption with 6000-km changes, liters/100 km; (d) change; (e) added; (f) average; (g) maximum; (h) oil consumption, liters/100 km; (i) oil consumption, % on fuel; (j) convoy No. 1202, Sevastopol'; (k) convoy No. 1126, Pskov'; (l) convoy No. 1141, Orel'; (m) "Livadiya" Fleet, Crimea.

by the centrifugal final filter. The inorganic part of total impurities does not exceed 0.2%.

Analysis to determine the additive content in the oil indicated that the barium content had dropped by 35-40% and zinc content by an average of 50%.

Thus, AS-8 oil with VNII MP-360 additive exhibits adequate antiwear, antioxidant and other operational properties and ensures dependable operation of ZIL-130 engines with the specified oil-change interval (every 4-8 thousand km) depending on operating conditions.

The results obtained on aging of the oil in the engine at 50°C (up to 200 thousand km) mileages confirm the correctness of the recommended oil-change interval.

#### Conclusions

1. AS-8 oil with VNII MP-360 additive, produced according to GOST 10541-63, has operational indicators that meet the requirements of a heavily loaded high-speed ZIL-130 engine.

2. In operation on AS-8 oil with additives, the parts of ZIL-130 engines show satisfactory wearability after up to 200 thousand km.

3. The oil-consumption [norm] for ZIL-130 engines can be lowered to 2.5% (on the fuel).

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| 1. Englin, K., <u>Porshneyye kol'tsa i smazka dvigatelya (Motor oils and engine lubrication)</u> , Gosoptekhizdat, 1959. |
| 2. Dzhordzhi, K., <u>Motornyye masla i smazka dvigatelya (Motor oils and engine lubrication)</u> , Mashgiz, 1963.        |

with consideration of the distinctive operating conditions of tractor diesels and the application of supercharging. The technical data of the engine are given below.

GOST designation	88441
Effective power $N_e$ , hp	210
Crankshaft speed $n$ , rev/min	1700
Cylinder bore, D, mm	130
Stroke S, mm	140
Number of cylinders 1	8
Average effective pressure $p_e$ , kg/cx	7.5
Average piston speed $C_p$ , m/s	7.95
Displacement $V_h$ , liters	14.86
Fuel consumption $G_f$ , g/(ehph)	170
Oil capacity V, liters	32
Oil consumption, % on fuel	not above 2

The bench tests of the oils consisted of two stages: - 240-hour short tests without changing the oil for preliminary evaluation and 800-hour extended tests with an oil change after 120 h.

The conditions of each test according to GOST 491-75 are given below.

Shaft speed, rev/min	1700	1300	1700	1700
Load, hp	none	180	210	none
Time, min	10	10	210	10

The indicators taken for evaluation were: quantity of deposits on piston-group parts, freedom of piston rings, piston-ring wear (based on weight loss), state of exhaust-valve seating surfaces, extent of clogging of centrifuge rotor chamber (HRS-reaction-type oil centrifuge), and the changes in the physico-chemical properties of the oil, especially alkalinity. The degree of piston-group fouling was also determined by the point system.

Diesel fuel containing 1% sulfur was used in all tests. The base oil was DS-11. A list of the oils given, short-term tests and the principal results are given in Table 1.

We may conclude from the data of Table 1 that all of the experimental additive oils ensure good piston-ring stability, but differ substantially as regards their detergent and dispersing properties.

The poorest results as regards piston-group and centrifuge rotor deposits were obtained in the tests of oil with 6 and 6 $\frac{1}{2}$  of VNII NP-360 additive.

Series I oils with the domestic additives ensure satisfactory cleanliness of the engine parts and appear to be substantially superior to VNII NP-360 additive.

#### TEST RESULTS FOR OILS WITH ADDITIVES USED IN THE YaMZ-238B ENGINE

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The Yaroslavl Engine Works has developed and put into production a new series of V-type, four-stroke diesels with outputs ranging from 180 to 520 hp, including supercharged versions. Use of the four-stroke cycle with compression ignition has made it possible to produce inexpensive engines with high wear resistance of their parts and good starting. The V-type cylinder arrangement is used as the basis for the small, compact powerplants. The engine designs provide for maximum interchangeability of parts and subassemblies. It was noted in elaboration of the designs and testing of the new engines on diesel oil with TSIATIM-339 [TSIATIM: Central Scientific Research Institute of Aviation Fuels and Lubricants] that this oil does not meet their requirements. It was therefore necessary to find oils for the four-stroke diesels, and especially for the supercharged models used in the K-700 tractors. Although satisfactory results had been obtained in preliminary tests of oil with the VNII NP-360 additive in the YaMZ-236 engine [YaMZ: Yaroslavl Engine Works], further experience in the operation of K-700 tractors with YaMZ-238NE engines showed that this oil does not ensure stable operation of the piston group.

An extensive battery of tests was carried out with the YaMZ-238NE engine (a four-stroke V-type eight-cylinder diesel with direct fuel injection) in order to establish quality requirements for the oils and select additives for the YaMZ engines

Table 1. Results of Short-Term Oil Tests

a. Oil design.	b. Producer and source	c. % chromium plated rings	d. % over-all scaling on piston and rings	e. % scale on piston and rings	f. % wear of top ring	g. deposits in RMTs over Monto 613;	h. % scale on skirt of piston and rings	i. % scale in slots of oil-control rings	j. % scale on piston and rings	k. % scale on piston skirt
i. JC-11+0.05% Chromate-433+	1. Септим 1	0.20	2.5	9.8	0.14	1250				
j. JC-11+0.4% Rimula 613	6% BHIM MI-360	0.35	3.0	14.2	0.04*	2080				
k. JC-11+0.005% INC-200A	8% BHIM MI-360	1.25	3.75	16.2	0.06*	2520				
l. JC-11+5% Chromate-433+	5% BHIM MI-360	0.20	2.5	6.4	0.37	1035				
m. JC-11+5% Chromate-433+	1. УНИТИМ-339+ИМСИ	0.70	18.0	4.1	0.22	740				
n. INC-200A	0.70	18.0	4.1	0.22	740					
o. JC-11+3% Бимини-2+1%	q. 3% Бантиш-2	1.05	2.0	7.68	0.15	1620				
p. JC-11+3% Бок-25+2% СР-3+	s. 5% БДК	0.67	2.0	7.9	0.18	2700				
q. JC-11+0.005% INC-200A	t. Септим 2	1.50	21.5	9.0	0.22	1520				
r. JC-11+5% Chromate-433+	u. 11% BHIM MI-370	1.50	24.0	9.4	0.23	1520				
s. JC-11+5% INC-200A	v. МАК+1% АХ+3% АФ-31+	0.85	13.5	3.6	0.42	570				
t. INC-200A	y. Паяза	1.0	4.9	4.1	0.25	600				

\*Chromium-plated rings.  
KEY: (a) products; (b) type designation of oil; (c) ash content of oil; (d) over-all scaling on piston, points; (e) amount of scale on piston and rings, g; (f) wear of top ring; (g) deposits in RMTs over test period; (h) DS-11 + 0.25% Santolube 493 + 6% Monto 613; (i) series 1; (j) DS-11 + 6% VNIIT NP-360 + 0.003% PMS-200A; (k) 6% VNIIT NP-360 + 0.003% PMS-200A; (l) DS-11 + 5% VNIIT NP-370 + 0.005% PMS-200A; (m) DS-11 + 5% VNIIT NP-370 + 0.005% PMS-200A; (n) DS-11 + 3% TSATIM-339 + 2% PMS-Ya + 0.003% PMS-200A; (o) TSATIM-339 + PMS-Ya; (p) DA-11 + 3% BashNII-3 + 1% PMS-Ya + 0.03% PMS-200A; (q) 3% BashNII-2; (r) D-11 + 5% BPK + 2% SB-3 + 0.005% PMS-200A; (s) 5% BPK; (t) DS-11 + 0.7% Santolube 493 + 5% Monto 613; (u) Series 2; (v) DS-11 + 1% VNIIT NP-370 + 4% PMS-Ya + 0.05% L-23k + 0.005% PMS-200A; (w) DA-11 + 3% ASK + 3% MASK + 1.2% DF-11 + 0.03% PMS-200A; (x) ASK + MASK; (y) Rimula.

Series 2 oils with Monsanto additives, like VNIIT NP-370, and other additives are no better than the Series 1 oils as regards formation of scale and varnish on the piston group and the amounts of ash deposits in the combustion chamber (exhaust valves, piston heads). The smallest amounts of deposits were observed for

Table 2. Fouling of Parts after Long-Term Tests

a. Oil design.	b. Producer and source	c. % chromium plated rings	d. % over-all scaling on piston and rings	e. % scale on piston and rings	f. % scale on piston skirt	g. % scale on skirt of piston and rings	h. % scale in slots of oil-control rings	i. % scale on piston and rings	j. % scale on piston skirt
k. 6% BHIM MI-360	k. 6% BHIM MI-360	0.00	59	34.5	6.6	33.2	5.7	37.8	100
l. 5% BHIM MI-360	l. 5% BHIM MI-360	0.00	57	35.7	6.7	33.5	6.4	21.7	750
m. 5% BHIM MI-360	m. 5% BHIM MI-360	0.00	72	15	5.6	33.0	6.4	35.5	750
n. 5% BHIM MI-360	n. 5% BHIM MI-360	0.00	33	21	4.7	31.7	6.4	31.7	650
o. 5% BHIM MI-360	o. 5% BHIM MI-360	0.00	35.0	24.5	5.6	31.5	6.4	32.4	1150
p. 5% BHIM MI-360	p. 5% BHIM MI-360	0.00	33	23	5.1	32.5	6.4	31.9	1150
q. 5% BHIM MI-360	q. 5% BHIM MI-360	0.00	35	15.1	4.9	31.5	6.4	31.9	1150

\*Oil-change interval 240 hours, elsewhere 120 hours.  
KEY: (a) type designation of oil; (b) test time, hours; (c) piston scaling, points; (d) total; (e) including; (f) on groove; (g) on skirt; (h) on drainare holes and in slots of oil-control rings; (i) amount of scale on piston and rings; (j) deposits in RMTs during test period; (k) 6% VNIIT NP-360; (l) 1% TSATIM-339 + PMS-Ya.

operation on low-ash series 2 oils with ASK and MASK additives and with the imported Rimula oil, and the piston-group components were cleaner by factors of 1.5-2 than after running on Series 1 oils. Ring wear was slight during the test period and practically the same for the various samples, except for the oils with ASK and MASK additives, which permitted increased wear of the first compression ring.

Oils that had produced encouraging results in the short-term tests were given long-term tests, along with oils containing 6 and 2% of the VNIIT NP-360 additive, which are currently recommended for K-700 tractors.

Table 2 presents piston-group fouling ratings from the 800-hour tests; they confirmed the data of the short tests in indicating unsatisfactory detergent properties for oils with ; and 8% VNIIT NP-360 additive.

As we see from Table 2, increasing the additive concentration from 6 to 8% does not reduce the amount of scale formed, as indicated by the approximately identical overall scaling ratings - 59 and 57 points, respectively. In the long-term tests and 39 and 37.5 points for the short tests. It was also noted that the top oil-control rings were less mobile and that the amount of deposits in the ring grooves and on the piston skirts was larger.

It was established in 800-hour tests on oils with VNIIT NP-360 additive that the drainage holes in the pistons and the

slots of the oil-control rings were 60-70% filled with carbon deposits, which might interfere with normal operation of the piston group if the engines were run any further. Also noted in operation on these oils was considerable fouling of the engine crankcase, clogging of the centrifuge rotor and the sections of the coarse filter. Table 3 gives the results of the long-term tests.

Table 3. Wear of Parts After Long-Term Tests

Parameter of performance (a)	b	c		d	e	f		g	h	i
		Number of rings	Wear, mm			Weight loss, mg	Wear, mm			
1% BHMK HI-360	800	0.205	0.31	14	36	12	10			
1% BHMK HI-360	800	0.172	0.29	14	43	10	11			
5% BHMK HI-360	800	0.46	0.73	35	25	5	5			
5% BHMK HI-360	1200	0.68	1.1	56	36	5	5			
5% BHMK HI-360	1000	0.57	0.765	49	33	8	8			
5% BHMK HI-360	800*	0.62	0.64	33	33	11	11			
KUHATM-360+PMSCY	800	0.12	1.02	60	46					

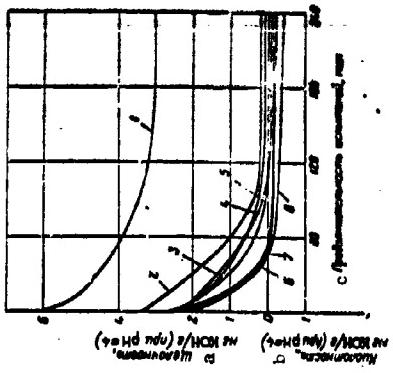
\*Oil change interval 240 hours.

KEY: (a) type designation of oil; (b) test time, hours; (c) weight loss; (d) first ring; (e) set of rings; (f) wear; (g) first ring, height dimensions; (h) first piston groove; (i) sleeve; (j) 6% VNII NP-360; (k) TSIAITM-339 + PMSCY.

Satisfactory results were obtained from tests of Series 1 oils with 5% of VNII NP-370 and with the TSIAITM-339 and PMSCY additives.

In 800-hour tests, oils with 5% VNII NP-370 preserved full piston-ring mobility and moderate piston-group point ratings; the drainage holes were practically clean and there were only small deposits in the centrifuge. The test time for this oil was increased to 2000 and then to 3000 h for the same engine, with the parts checked for cleanliness and then cleaned after each test stage. With increasing engine running time, we note some increase in the fouling of parts as the piston group wears. Then 800-hour tests were conducted with a 240-h oil-change interval, with the result that scaling on the piston group increased to 39 points, while the amount of sludge increased from 8.6 to 12.9 g. Even in this case, however, the parts were cleaner than after operation on oil with VNII NP-360 additive.

Oil with 6 and 8% of the VNII NP-360 additive indicated superior antiwear properties in the long-term tests. Some increase in the wear figures was noted after operation on oil with the additive TSIAITM-339 + PMSCY (see Table 3).



Change in alkalinity of oil:

- 1) ASK + MASH; 2) 5% VNII NP-370;
  - 3) Series 1; 4) TSIAITM-339 + PMSCY;
  - 5) BashMII-2; 6) BPK + SB-3; 7) 6% VNII NP-360.
- KEY: (a) alkalinity, mg of KOH/g  
(at pH = 4); (b) acidity, mg of KOH/g  
(at pH = 4).

Oil quality (apart from detergent and antiwear properties) was also evaluated from the chamber-surface condition of the exhaust valves, which were made from EI-69 steel (both without facing of the chamber and with a VZK-alloy facing). It was found after the tests that the built-up valve chambers were in better condition than the unfaced chambers for all of the oils.

It should be noted that formation of a glasslike film on the valves (both faced and unfaced types) had been noted previously in a number of cases. The gases are usually puffed into cracks and pits in this film, occasionally burning valves.

This glasslike film on the valves was not observed in our tests of oil with VNII NP-360 additive.

The problem of glass-film formation on the exhaust-valve chambers in operation on oils with certain additives requires special study.

Particular attention was devoted to the change in the oil's neutralizing properties in the oils during these tests.

For this purpose, the alkalinity of the crankcase oil was determined during the testing process. The figure shows the analytical results. Alkalinity drops off most sharply in the oil with 6% of VNII NP-360. With this specimen, strong acids were detected even after 60 hours of operation, and their content subsequently increased. Increasing the additive concentration to 8% had little influence on the neutralizing-property change.

Oil with 5% of the additive VNII NP-370 has the largest alkalinity reserve. Oil with the combined additives TSIATIM-339 + PMS Ya has almost the same alkalinity. The alkaline properties are retained in these oils until they are drained; in other oils of Series 1, high acidity is found after as little as 120 hours of operation. Series 2 oils retained their alkaline properties to the end of the test.

The figure shows that the alkalinity of the oil drops most rapidly during the first few hours of operation, and that this drop is more abrupt in the oils with the VNII NP-360 and BPK + SB-3 additives. Oil alkalinity then stabilizes as a result of addition of fresh oil and filtering out of the oxidation products. The data plotted in the figure are in good agreement with the results of engine-parts inspection after the tests.

#### Conclusions

1. It was established by the tests that the best oil for the YaMZh-238NB engine is the Series 1 oil with 5% VNII NP-370, 2% PMS Ya, 0.5% LZ-23k and 0.005% PMS-200A. This oil was recommended for field testing.
2. Oil with the TSIATIM-339 + PMS Ya additive, whose properties resemble those of the Series 1 oils, is also recommended for use testing.
3. Rather heavy deposits are formed on the piston group, in the centrifuge rotor, and in the engine crankcase when the engine is run on oil with the VNII NP-360 additive; this additive does not meet the requirements for dependable operation of the engine on sulfur-containing fuels.
4. Operating experience and the results of testing show that different batches of oil with the VNII NP-370 differ in quality. The physicochemical indices stated in the GOST do not give a complete picture of the motor properties of the oil. It is therefore necessary to formulate and introduce into the GOST quick oil-property motor-rating methods, so that the refineries will be able to indicate their results on the tags accompanying the oil.

#### **EFFECTIVENESS OF USING ADDITIVES IN HYDRAULIC-COUPLING OILS FOR PASSENGER AUTOMOBILES**

O.S. Obieukhova and V.N. Kryucheshnikov

The operating conditions of the oil used in rotor-vehicle hydraulic transmissions - automatic transmissions and torque converters, power-steering gear, etc. - are characterized by: 1) broad temperature range (from -60°C for vehicles used in the northern regions to +130°C); 2) high specific pressures; small clearances (especially in automatic transmission controls); by the presence of parts containing substantial amounts of copper (up to 60% in sintered brakebands); by contact with rubber gaskets and hoses.

None of the mineral oils can guarantee dependable long-term service from these units, since these oils cause increased wear and formation of deposits that interfere with operation of the automatic controls and other assemblies.

VNII NP-1 oil, which is made by using polyisobutylene to thicken a light deep-refined and deparaffinized oil with DP-1, phenyl- $\alpha$ -naphthylamine, and VNI-5K additives, is produced at the present time for passenger-car automatic transmission. These additives are used to improve the viscosity, antiwear, anticorrosion, antifoaming, and antioxidant properties. The quality indicators of this oil are given in Table 1.

User experience with VNII NP-1 oil has indicated that its quality must be improved, since the test constituents of the oil had a pour point of minus 40°C, while the standard specifies minus 35°C and commercial consignments pour at minus 32°C. Use

Table 1. Oil Properties

a Measure	b VNII MP-1 100°C	c VNII MP-1 100°C	d VNII MP-1 100°C
e Boiling, °C	260	260	260
f Pour, °C	-30	-32	-32
g Copper-plate corrosion, "C"	-35	-39	-46
h Pour point, °C	30	30	30
i Viscosity at 100°C, mm²/sec	1.15	0.03	0.03
j Viscosity at 40°C, mm²/sec	74	73	73
k Viscosity at 100°C, mm²/sec	24	23	23
l Viscosity at 100°C, mm²/sec	±20	+45	-18

KEY: (a) indicator; (b) VNII NP-1 GOST 10660-63; (c) thickened base oil of oil VNII NP-1 + MASK + DP-11; (d) AU spindle oil + MASK + DP-11; (e) viscosity, cst; (f) at: (g) pour point, °C; (h) copper-plate corrosion in 3 hours at 100°C; (i) temper color; (j) none; (k) thermal oxidation stability in DK-2 machine; (l) sediment, %; (m) acid number, mg of KOH/g of oil; (n) reaction, %; (o) weakly alkaline; (p) neutral; (q) tests on ChShM-3; (r) seizing load, kg; (s) welding load, kg; (t) OPI [generalized wear index]; (u) swelling of type 78 rubber in 72 hours at 130°C.

of the oil is limited by its high viscosity at low temperatures.

The oil has excessive corrosive aggressiveness toward copper. Copper-plate tests at the operating temperatures of the oil (10-130°C) indicate the formation of a black, loosely attached film that accelerates part wear. The oil has inadequate antiwear stability. This makes it necessary to change it at least every 26 thousand kilometers.

The metric value obtained by GOST 10660-63 for rating the stability of oil is 10. The VNII [All-Union Heat Institute] machine does not characterize the behavior of the oil in use; neither does holding the oil at 140°C for 24 hours. In both cases because of the mild oxidation conditions.

The NAMI [Central Automotive-Engine Scientific Research Institute] DK-2 machine was used for stability evaluation of the oils; VNII NP-1 and several Type A (General

Motors) oils developed abroad for the same application were tested in it at various temperatures.

It was established that sediment forms after 21 hours (at 160°C oxidation of VNII NP-1 oil), and that it amounts to 1.6% after 50 hours, while sediment forms in the imported oils only after 177-182 hours.

A copper plate with the same dimensions as the lead plate used in determining the corrosive aggressiveness of oils in the NAMI DK-2 machine was used as a catalyst. This made it possible to shorten the testing time significantly (by 30-40%) and to test the influence of the oil on the copper plate. The following method of evaluating the stability of oils used in hydraulic systems was selected on the basis of a series of tests: after oxidation in the NAMI DK-2 device for 60 hours at 160°C in the presence of the copper plate, the oil may not form sediment even on subsequent solution in ten times the amount of isooctane (or petroleum ether) and storage of the solution for 24 hours. Formation of a black film that cannot be rubbed off the copper plate is unacceptable.

Additives that confer improved stability on VNII MP-1 oil were selected by the method indicated.

The best results were obtained with VNII MP-1 oil to which the following additives were introduced: MASK (4-4.5% with a calcium content of 0.15-0.18%), DF-11 (2-2.2% at a zinc content of 0.1%) and PMS-200A (0.005%) as an antifoaming additive.

The laboratory tests showed that the experimental oil has high antioxidant stability and good antisear and anticorrosion properties (see Table 1). It was also established that MASK additive made from a heavy raw material acts as a depressor and lowers the pour point of VNII NP-1 oil to minus 49°C. When MASK additive made from light raw materials is used, the pour point of the oil can be lowered by adding 0.3-0.5% of polymethacrylate D.

VNII MP-1 oil with the new additive combination was tested in the fluid transmissions of ZIL-111 passenger cars. The transmission parts were measured with the micrometer at the beginning of the tests and after 50 thousand kilometers. Oil samples were taken for control purposes at 25-thousand-km intervals. The oil was not changed until the vehicles had accumulated 55 and 100 thousand km. The test results were compared with data obtained with VNII NP-1 oil and imported Shell Dorax T-6, which were changed every 25 thousand km.

The results of use tests with the experimental oil were evaluated on the basis of deposits formed on transmission parts and in the transmission sump, transmission trout le, the wear of its parts, the behavior of the rubber packing components, and the change in oil quality during operation.

Use of the test oil without changes for 55 and 100 thousand km kept the transmissions running normally. When the units were disassembled, no oil-oxidation products could be detected in the form of gummy deposits, varnishes, etc., either in the automatic control unit (control panel), reduction-valve panel, housing of centrifugal spool) or on other hydraulic-transmission or torque-converter parts. There was 5-6 g of sludge in the bottom of the case after 50 thousand km, and it consisted basically of inorganic impurities.

When the transmissions were run with VNII NP-1 oil, there were ~20 g of sump deposits at 25 thousand km. Sludging was also observed in the passages of the panel case and in the reduction valves, in some cases causing them to stick.

No parts of the hydraulic transmissions run on the test oil showed deep scoring or temper colors. For the most part, the parts had worn by 0.01-0.015 mm after 93 thousand km; only on one part did the wear amount to 0.08 mm (as against a maximum acceptable wear of 0.2-0.3 mm).

The clutch is the basic element in determining the dependability and service life of a hydraulic transmission. In operation on the test oil, the wear of the sintered-metal clutch disks was practically nil, and no dark films formed on their surfaces. The test results indicated that the test oil is on a par as regards antiwear properties with the imported Shell Donax T-6. When VNII NP-1 oil was used, the sintered parts showed more rapid wear. Analysis of oil samples taken during the tests showed that the viscosity level dropped somewhat in both the commercial VNII NP-1 oil and in the test oil (by 3-4 cst at 50°C) as a result of destruction of the viscosity additive.

The oxidation-product content of the test oil (the organic part of the total impurities) was insignificant, and did not exceed 0.02-0.03% over the entire useful life of the oil (93 thousand km), while it was 0.54%, including 0.32% of asphaltenes in the VNII NP-1 oil and the

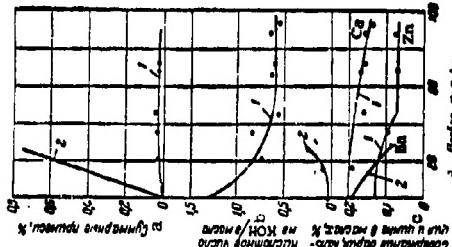


Fig. 1. Changes in additive-component contents, acid number and total impurities in oil as a function of engine mileage. 1) test oil; 2) VNII NP-1 oil.  
KEY: (a) total impurities, %; (b) acid number, mg KOH/g of oil; (c) barium, calcium and zinc contents in oil, g; (d) mileage, thousands of km.

inorganic part of the impurities (0.2%) contains a considerable amount of spent DF-1 additive. For this reason, the acid number of the oil decreased considerably (Fig. 1).

The concentration change of the MASK and DP-11 additives in the test oil during operation was insignificant (see Fig. 1). The following conclusions may be drawn from the test results.

Introduction of the MASK, DP-11 and PGS-200N additives into the thickened VNII NP-1 oil base produced passenger-car automatic-transmission oils with good antiwear, anticorrosion, and antioxidant properties. The depressor properties of the oil can be improved by introducing 0.3-0.5% of polymethylacrylate D.

The high stability of this oil makes it possible to increase the change interval substantially and reduce operating consumption. Even with changes at 50 thousand km, oil consumption is reduced by 40-45% by comparison with the commercial VNIK NP-1 oil.

**Power-steering oil.** The oil used in automatic transmissions and torque converters is usually also used in power-steering systems. Good viscosity-temperature properties of this oil are especially important during starting of the engine and steering-booster pump. VNII NP-1 oil is fully adequate for passenger automobiles, but its viscosity-temperature properties have been found unsatisfactory for trucks and buses, which are generally parked outdoors.

It was established by special tests run on 2NL-130 vehicles that at air and pump-oil temperatures of minus 55°C, the oil temperature is raised to minus 40°C during the prestart warmup of the engine, and then reaches 0°C within 1-2 min after the pump has started. Under these conditions, normal operation of the pump can be ensured by use of an oil with a minus 40°C viscosity not above 15-18 thousand cst and a pour point below minus 45°C.

It should be noted that use of ACM and AMG-40 light commercial oils, which have good viscosity-temperature properties, results in increased wear of parts.

The viscosity of the oil is lowered when VNIK NP-1 oil is used in the steering hydraulic booster; after 4 thousand km, the 50°C viscosity has fallen from 26 to 16-18 cst. This substantial viscosity drop is explained by destruction of the viscosity additive, which results both from the high running temperature of the oil (up to 130°C) and the incessant stirring (since the pump delivers 10 liters/min at 600 rev/min and the excess of oil is directed through the transfer valve when crankshaft speed changes).

Thus, acquisition of an oil with good operational properties requires: a base with good viscosity-temperature properties; a viscosity additive with high thermal and mechanical stability; an antwear additive that ensures long service life and dependable operation of the power-steering gear in all of the country's climate zones; an antioxidant to provide a long oil-service life, and an antifoaming additive.

In developing the oil, we took as the base AMG-10 oil with various viscosity additives (vinipol, polyisobutylene, polymethacrylate) and type AY spindle oil from Zhirkovsk petroleum. The antiwear and antioxidant additives for these oils were first tested in various combinations on a friction machine and then checked on the DK-2 machine in order to obtain test specimens whose antiwear properties and stability would meet the specifications for automatic-transmission oils.

The following additive combinations were chosen: MASK (4%), and EP-11 (2.3%); chlorinated paraffin (5%) and MB-11 (1%).

**Running In:** pump load 125 cycles in 1 min; oil pressure drop across system under various loads:  $P_1 = 10-20 \text{ kg/cm}^2$  for 10 hours, oil temperature 115-125°C;  $P_2 = 10-45 \text{ kg/cm}^2$  for 10 hours, oil temperature 115-125°C. Pump shaft speed 3000 rev/min.

The normal operating regime of the pump differs only in having a higher pressure ( $10-55 \text{ kg/cm}^2$ ).

The quality of the oils was evaluated from the change in their physicochemical constants and from the wear of the piston rod, characterized by the increase in its smallest diameter and its loss of weight (readings taken every 50 hours). The tests lasted 150-200 hours. Each oil specimen was tested in four or five pumps.

#### TEST RESULTS

The results of the pumpings at Figs. 2 and 3, from which we have  
See page 39 for footnote.

The results of the pump stator wear readings are plotted in FIGS. 2 and 3, from which we see that:

The bench tests showed that the viscosity additives undergo considerable destruction. Figure 1 shows the changes in

stator wear reaches its limit (1.25 mm) after 50 hours of running on AMG-10 commercial oil;

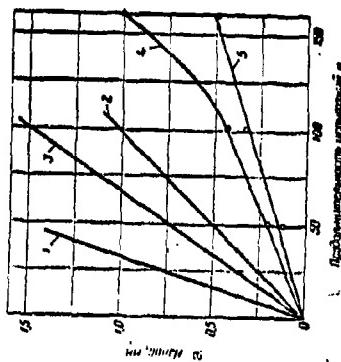


Fig. 2. Pump stator wear on various commercial oils. (1) AMG-10 with polymethylacrylate, chlorinated paraffin, and MB-1; (2) AMG-10 with polyisobutylene, chlorinated paraffin, and MB-1; (3) AU spindle oil; (4) AU spindle oil; (5) AU spindle oil with chlorinated paraffin and MB-1.

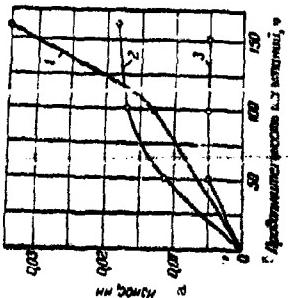


FIG. 3. Pump stator wear on various oils. (1) MW-10 with polyisobutylene, MASH; and DR-11; (2) VNIIT NP-1; (3) AU (spindle with MASH and DR-11).

on introduction of the MASK and DP-11 additives into these oils, wear is reduced substantially, especially in the case of bindable AU, where the wear ranges from hours of testing;

There is practically no stator wear when the injected Shell Donax T-6 oil is used. The best results as regards antiwear properties were obtained with the domestic MASK and DPM-11 additives.

Viscosity Properties

The bench tests showed that the viscosity additives undergo considerable destruction. Figure 1 shows the changes in

### Viscosity during the tests.

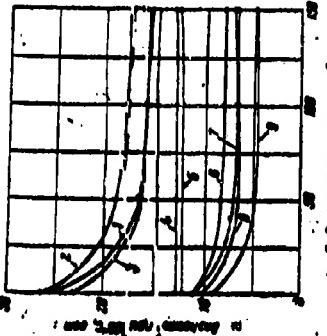


FIG. 4. Change of oil viscosity as a function of test time. 1) commercial VNII MP-1; 2) VNII MP-1 with polyisobutylene, MASK and DP-11; 3) import type A; 4) spindle AU; 5) spindle AU with MASK and DP-11; 6) AMG-10 with Vinipol, chlorinated paraffin and MB-1; 7) AMG-10 with polyisobutylene, chlorinated paraffin and MB-1; 8) AMG-10 with polyisobutylene, MASK and DP-11; 9) AMG-10 with polymethacrylate, chlorinated paraffin and MB-1.

KEY: (a) 50°C viscosity, cst; (b) test time, hours.

The total impurity content is insignificant in all oils, ranging from 0.01 to 0.02%, with asphaltene completely absent from the filtered oil. The acid number of AU spindle oil rises from 0.04 to 0.15 mg of KOH/g, but in oils containing the MASK and DP-11 additives, it declines as the additives are depleted, from 1.26 to 1.11 mg of KOH/g. However, the color of the test oil did not change during the tests, indicating that it has good stability to oxidation.

The bench tests made it possible to recommend spindle oil AU with the MASK, DP-11, and PMS-200A additives for road testing in the country's various climate zones.

### Conclusions

1. Introduction of the MASK (~4%), DF-11 (~2.5%) and PMS-200A (0.005%) additives into oil improves their antiwear, antioxidant, anticorrosion, and antifoaming properties considerably, with the result that the dependability and service lives of automotive automatic transmissions and power-steering pumps are substantially improved.
2. The high antioxidation stability of oil with these additives makes it possible to use it 2-3 times longer and reduces operational oil consumption.

The greatest viscosity loss is observed when polymethacrylate (51.6%) and polyisobutylene (32.40%) are used; the viscosity decrease is smaller with Vinipol (19.4%). The viscosity level of the imported oil also falls by 30%.

Characteristically, the viscosity drops during the first few hours of operation, with some stabilization observed after 50 hours.

The viscosity of AU spindle oil rises slightly (by 0.39 cst at 50°C) as a result of formation of asphalt tars. The stability of the oil is raised by introduction of additives, so that its viscosity remains practically constant (0.09-0.2 cst at 50°C after 150 hours of testing).

These tests showed that commercial viscosity additives with molecular weights not below 10,000 should not be used in power-steering oils. Distillate oils such as spindle AU, which have lower viscosity levels at the working temperatures, lubricate the rubbing surfaces satisfactorily in the presence of antiwear additives.

The antioxidation stabilities of the oils cannot be determined conclusively from the results of bench tests because of their short duration (150-200 hours). The total impurity content is insignificant in all oils, ranging from 0.01 to 0.02%, with asphaltene completely absent from the filtered oil. The acid number of AU spindle oil rises from 0.04 to 0.15 mg of KOH/g, but in oils containing the MASK and DP-11 additives, it declines as the additives are depleted, from 1.26 to 1.11 mg of KOH/g. However, the color of the test oil did not change during the tests, indicating that it has good stability to oxidation.

## COMBINATION OF ASH-FORMING AND POLYMERIC ASH-FREE ADDITIVES IN MOTOR OILS AS A WAY TO FURTHER QUALITY IMPROVEMENT

<sup>1</sup>M-1 additive (*4*,*4*-methylene-bis-*2*,*6*-tert-isobutyl-*1*-phenol).

Ecclesiastes

Manu-  
script

<sup>1</sup>MP-1 additive (*4,4'-methylene-bis-2,6-tert-isobutyl-phenol*).

FORMATION OF ASH-FORMING AND POLYMERIC ASH-FREE ADDITIVES  
MOTOR OILS AS A WAY TO FURTHER QUALITY IMPROVEMENT  
Ye.G. Semeido, I.A. Traktovenko, N.V. Shevagolev

As we know, only metal-containing organic compounds (ashing additives), and especially calcium and barium alkylphenolates and their sulfur- and phosphorus-containing derivatives, are used as detergent, dispersing, anticorrosion and antioxidant additives to motor oils produced by the domestic petroleum industry. Sulfonates of alkaline-earth metals and zinc dialkyl-dithiophosphates are also used.

Depending on application, motor oils may contain 6-15% more of these ashing additives. However, when engines are run on oils containing large amounts of ashing additives, their rubbing parts, and especially their cylinder-piston group, wear rapidly. This wear is a consequence of carbonization of the oil films in hot areas of the engines, with formation of the cinders of the metals.

The "hotter" and more powerful the engine, the more rapid is the depletion of the ashing additives in its motor oil. To compensate for depletion, larger amounts of the additives are introduced into the oils. Depletion of the ashing additives is also influenced by cold engine operation under city traffic conditions. When the hydrophilic properties of the ashing additives (salts of alkaline-earth metals) are manifested most strongly, i.e., formation of low-temperature deposits is intensified.

Thus, together with their good detergent and dispersing properties, ashing additives have important disadvantages that make it necessary to reduce the content of these additives in engine oils.

The necessary detergent, dispersing, and antiwear properties can be obtained in motor oils with minimum contents of ashing additives if nitrogen-containing polymer (nonash) additives are introduced into the oils.

The nonash additives belong to the class of polar polymers and are copolymers of methacrylic acid alkyl esters with certain nitrogen-containing monomers, such as methylvinylpyrrolidine, diethyldiaminoethylmethacrylate, etc.

In the research whose results are reported below, we used VNM nonash additives prepared by V.G. Telegin and Ye.V. Lazareva [1].

To determine the effectiveness of the nonash additives in combination with ashing types, a motor oil containing these additives was tested on a YaMZ-236 diesel engine. The test results were compared with the results of tests made with the same engine under similar conditions using motor oils with only the ashing additives. Each sample was tested for 150 hours.

The detergent, dispersing and antiwear properties of the additives are evaluated on the basis of:

- the quantity and type of deposits on the engine parts and oil filters;
- piston-ring freedom;
- the wear of engine parts and their condition;
- the change in the physicochemical properties of the drained oil.

It has been established by previous studies and tests [2] that 5% of ashing additives (5% SB-3, 2% DF-11) can be used in motor oils for type YaMZ-236 diesels and type ZIL-130 V-type carburetor engines. In motor oils made from eastern petroleums and used in diesels, this additive content gives only satisfactory results.

To ascertain the possibility of obtaining a sharp increase in the quality of oil containing the acceptable amount of ashing additive by adding the VNM nonash additive to it, we tested oil A, which was type ASZp-10 containing 3% SB-3, 2% DF-11 and 5% VN-18 (Table 1).

The results were compared with the results of tests made with oils having high ashing-additive contents: oil B, which was

Table 1. Results of Analysis for Physicochemical Properties

a Series	b Viscosity at 100°C, cSt	c Ozone resistance at 25°C, min	d Coking capacity at 100°C, %	e Ash content, %	f Acid number, mg of KOH/g	g Temperature, °C	h Flesh point, °C	i Pour point, °C	j V
A	1.05	3.69	0.339	0.41	0.654	1230	115	100	-40
B	10.38	0.23	0.329	0.43	0.132	1230	115	105	-35
C	10.30	0.25	0.325	0.43	0.132	1230	115	105	-35
D	10.21	0.37	0.326	0.41	0.021	1230	115	105	-35

KEY: (a) oil; (b) viscosity at 100°C, cSt; (c) ratio; (d) ash content, %; (e) coking capacity, %; (f) acid number, mg of KOH/g; (g) temperature, °C; (h) flesh point; (i) pour point; (j) B; (k) V; (l) E.

Table 2. Deposits on Engine Parts

a Series	b Coke-forming deposits at 25°C, %	c Ash-forming deposits at 25°C, %	d Varnish deposits at 25°C, %	e Ash content on parts, %	f Ozone resistance on parts, min	g Ash content on coarse filter, %	h Ash content on fine filter, %	i Varnish content on coarse filter, %	j V
A	50-55	30-70	10-15	23.3	100	55	55	55	-42
B	35-45	20-45	25-30	22.2	100	55	55	55	-42
C	30-35	20-30	20-25	24.1	100	55	55	55	-42
D	1	1	1	1	1	1	1	1	1

KEY: (a) oil; (b) condition of pistons; (c) % of piston-shirt area coated by varnish; (d) % of piston head area covered by scale; (e) PZV result, points; (f) deposits on parts, %; (g) total amount of scale; (h) deposits in centrifugal filter; (i) deposits in coarse filter; (j) P; (k) V; (l) E.

type ASZp-10, 4% of ash-forming additives (7% SK-11 and 2% DF-11), and 1% V, which was type DSp-11 with 6% of VNIIT No. 360, as well as a standard (E) oil of type DSp-11, 4th 4% Monto 613 and 0.25% Santolube 493, the latter corresponding to Series I.

It is known that the serviceability of an engine is largely determined by the deposits on its parts. They are responsible for most cases of engine trouble. Table 2 presents data on the deposit-forming properties of the oils.

Piston-ring freedom is judged from the number of rings that can be shifted with little effort (there were no stuck rings in any of the tests); there were two such rings in the tests of oil A and three each in the tests of oils B, V and E.

Table 3. Test Results

Series	b. Engine service test results											
	a	c. Crank bearings, mm	d	e. Impregnation factor, %	f	g	h	i	m	n	o	
A	6.9	7.0	7.5	0.10	0.05	0.03	0.01	0.05	0.03	0.01	0.05	
B	6.9	7.5	8.0	0.13	0.05	0.04	0.01	0.05	0.02	0.02	0.05	
C	6.9	8.5	9.5	0.11	0.04	0.03	0.01	0.04	0.03	0.02	0.07	
D	6.9	8.5	7.0	0.13	0.06	0.04	0.01	0.06	0.02	0.02	0.05	

KEY: (a) oil; (b) engine-part wear; (c) cylinder liners, parallel to crankshaft axis; (d) average; (e) piston rings, E; (f) first compression ring; (g) second compression ring; (h) third compression ring; (i) first oil-control ring; (l) second oil-control ring; (m) connecting-rod inserts, E; (n) top; (p) bottom; (q) V; (r) E.

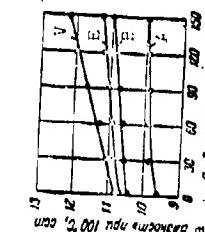


Fig. 1. Change in viscosity of oils during running in YaMZ-236 engine.  
KEY: (a) 100°C viscosity, cst; (b) oil service time, hours.

The wear and condition of the engine parts were approximately the same after tests with each of the above oils. There were minor differences in piston-ring and connecting-rod-insert wear (Table 3). The surface of the oil pan was clean after the tests of oils A and E, but it was covered with gummy deposits in the other cases.

Changes in the physicochemical properties of the used oils indicate high stability for oils A (with the nonashing additive)

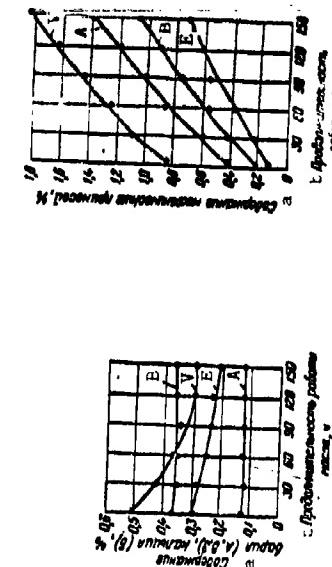


Fig. 2. Change in ash content of oil during running in YaMZ-236 engine.  
KEY: (a) ash, %; (b) oil service time, hours.

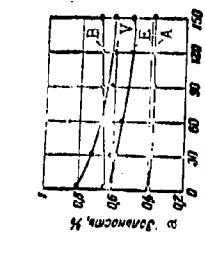


Fig. 3. Change in metal content of ash additives in oils during running in YaMZ-236 engine.  
KEY: (a) contents of barium (A, V, E) and calcium (B); (b) oil service time, hours.

and E (the reference oil) by comparison with oils V, which contained 6% of VNI NP-360 (Figs. 1-3). But even at increased concentration (6%) in ordinary oil, VNI NP-360 additive does not fully guarantee Series 1 quality.

Fig. 4. Results of Tests on GAZ-51 Engine

a	c. Special service engine test results											
	b	c. Oil consumption, liters/hour	d	e	f	g	h	i	j	k	l	m
A	0.9	0.004	0.002	0.018	0.017	0.014	0.007	4	5	1	1	1
B	1.5	0.005	0.003	0.023	0.024	0.019	0.010	3.5	3.5	1	1	1

\*Oil specimen with 3.5% VNI NP-360 additive tested.  
\*\*The denominator gives the result for oil with 6% VNI NP-360.  
KEY: (a) oil; (b) average diametral cylinder wear, micrometers; (c) average piston-ring wear, E; (d) compression rings; (e) oil-control rings; (f) average insert wear, E; (g) top; (h) bottom; (i) amount of deposits, g; (j) in crankcase; (k) in valve cover; (l) total; (m) V; (n) E.

The detergent and especially the dispersing properties of this additive are clearly inadequate to ensure normal diesel performance. Thus, FIG. 2 and Table 2 indicate that oil V with the VNII MP-360 additive contained 1.8% of mechanical impurities by the end of the tests, there were 820 F. of sludge in the centrifugal filter, while the corresponding figures for oil E (Series 1) were 0.7% and 365 g., for oil A (with the nonashing additive) 1.4%, and 190 g., and for oil B 1.1% and 510 g.

The piston-wear rating after the tests was 3-3.5 points for oil V and 2-2.5 and 1-1.5 points, respectively, for oils E and A.

A consequence of the inadequate dispersing properties of VNII MP-360 additive is a relatively sharp increase in oil viscosity during work (see FIG. 1). The fast depletion of this additive (see FIGS. 2 and 3) is also partly responsible for its decreased effectiveness.

Comparative 28-hour cold tests run on the GAZ-51 engine with oils having ashing additives and a mixture of ashing and non-ashing types indicated a relatively sharp increase in oil viscosity during work (see FIG. 1). The fast depletion of this additive (see FIGS. 2 and 3) is also partly responsible for its decreased effectiveness.

Inspecting the above data, we may conclude that combining ashing and nonash additives in a motor oil endows it with high operational properties, while the ashing additives taken alone improve one indicator at the expense of others.

A nonash additive in the combination of additives introduced into a motor oil improves its viscosity-temperature properties (VI, viscosity at 0°C, etc.).

The quality margin of oil A indicates the possibility of reducing the proportion of ashing additives in the oil when they are used in combination with the nonash additive. This is also confirmed by the good results of the 100-hour test run in the YaMZ-236 engine on a thickened oil containing the type VN nonash additive, 2.5% or SB-3, and 1% or LAMI-317.

In the production of ordinary motor oils containing ashing and ash-free additives (in various combinations), it is necessary, in view of the thickening tendency of the latter, to take initial distillate or residual oils with somewhat lower viscosity levels than when the ashing additives are used alone.

#### References

1. Soviet Patent 160,252, 8 February 1963. Byull. Izobr., No. 3 (1964).

BLOWING OUT INTO THE Cylinders, EXCLUDING PRESSURE, BY  
Other Engines and Main, Then at the Selected Level.

It must be possible to vary the rank and temperature of the fuel in the selected range of the calorimeter, and to regulate the electric heater walls.

The cylinder pressure is to be adjusted to the required level, and the temperature of the cylinder walls is to be maintained as nearly as possible at constant level, by means of electric heaters, which are connected to the cylinder walls.

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### ELECTION OF OPTIMUM ADDITIVE COMBINATIONS FOR MOTOR OILS ON A SEVEN CYLINDER, SINGLE CYLINDER AIR-COOLED ENGINE

#### 1. TESTS ON THE ENGINE

Several and different additive combinations were tested on the engine, and the results do not correspond to each other. After the trials, the conclusions drawn were that there was no substantial improvement, and often degradation, in the best results. Therefore, the additives are carried out in the test, and the results are referred to the samples of the best engines and the best additives of the sample.

In view of this, work has been done toward the creation of a special apparatus for testing oils and additives for use in aircraft engines and development of a motor rating method for aircraft engines and their properties at high temperatures.

This equipment was designed with consideration of the following requirements.

It is necessary that the test engine be sensitive to thermal stresses and other stresses, have a degree of thermal stressing and other properties, aid in simulating various modern engines, permit operation without breakdown on a broad quality range of oils, and permit differential evaluation of these oils.

The design of the engine should permit regulation of the oil and fuel consumption, the temperature of the oil,

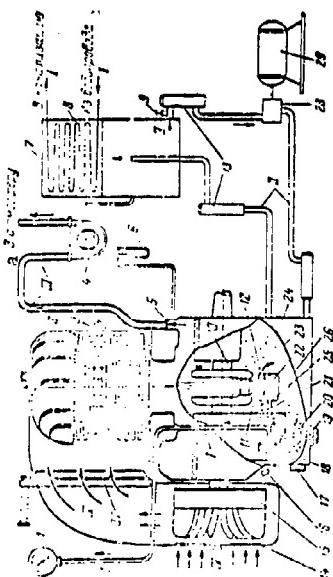


FIG. 1. Diagram of lubricating, cooling and ventilation system of crankcase in KMA-1 installation. 1) pressure gauge (0-1 kg/cm<sup>2</sup>); 2) device for regular rate of flow into engine crankcase; 3, 9) chrome-alum thermocouples; 4) type GK gas meter; 5) oil-filler pipe; 6) carbon manometer; 7) T-9 centisentri; 8) cooling coil; 10) holes for lubrication of connecting rod bearing; 11) oiling hole; 12) drain hole; 13) flywheel can; 14) regulating baffles; 15) temperature controller; 16) oil pump; 17) drain plug; 18) reducing valve; 19) oil pan; 20) oil pickup; 21) reducing valve; 22) oil tank; 23) circulation of gases; 24) glycol (ethylene glycol); 25) supplementary louver-type baffles; 26) KSh-2A gear pump; 27) AOL 32-4 electric motor ( $N = 1 \text{ kW}$ ). Lines: I) motion of oil under pressure; II) circulation of ethylene glycol or glycerine; III) removal of gases blown into engine crankcase; IV) motion of cooling air. KEY: (a) to atmosphere; (b) to sewerage; (c) tap water.

The quantity of blown-by gaseous is measured with the type GK gas meter ...

6. The temperature state of the engine is monitored by the use of thermocouples and a type EPP-09 recording electronic potentiometer.

An electric brake is provided by a type EL 1/4 induction motor (power 5.8 kW,  $n = 1450 \text{ rev/min}$ ).

The oil-testing unit thus devised with its single-cylinder carburetor engine was given the designation KMA-1.

The following test conditions were adopted as a result of preliminary results.

TEST TIME, HOUR	10
Effective effort, rev./min	360.1
Shaft speed, rev./min	204.25
Hour fuel consumption, kg/h	0.750.01
Spark advance, degrees	14.5
Cylinders	1.39+2
Oil in crankcase	1.45+1
Per-hour oil consumption, l/h	10.71
Avg. of oil tested, kg	0.71

To determine the depletion of the oils and additives, we conducted tests lasting up to 40 hours with variation of indicators after 10, 20, 30 and 40 hours without oil changing, but with 120 g added every 10 hours.

Oil quality was rated on the basis of the following indications:

the motor index  $I_m$  in points, reproducibility  $\pm 10\%$ ,

the deposit index  $I_d$  in points, reproducibility  $\pm 10\%$ ,

piston-ring freedom,  $P_k$ , in points, reproducibility  $\pm 30\%$ .

The scale against which ring mobility was evaluated is given in Table 1.

Table 1. Scale of Ring-Mobility Rating

Condition name	Open scale	Closed scale	Open scale
Quicke ring mobility	0	3	1
Chloroethylene saturation	1	2	2
Chloroethylene	2	3	3
Chloroethylene chloroform mixture	3	4	4
Chloroethylene	4	5	5
Hydrocarbon	5	6	6
Hydrocarbon	6	7	7
Hydrocarbon	7	8	8
Hydrocarbon	8	9	9
Hydrocarbon	9	10	10

KEY: (a) condition of rings; (b) rating, points; (c) clean, before testing; (d) drop freely with metallic sound; (e) drop freely; (f) drop freely with dull sound; (g) inert; (h) stuck; (i) slightly; (j) partially; (k) completely; (l) turned.

The arithmetic mean value was used in the calculation to determine the mobility rating for each ring.

The color of the deposits in the piston-ring zone ("yu") and on the piston skirt ("yu") was determined from the VIII-1961 Main Scientific Research Institute of Oil and Gas Processing

and Production of Artificial Liquid Fuels] color scale. The scale for rating varnish deposits on aluminum pistons consists of 11 color standards, ranging from light gray, which corresponds to the color of the clean piston and is rated zero, to black, which is represented by 10 points. The colors of the varnish deposits on the piston skirt and in the ring zone were rated separately.

To evaluate the deposits, the ring zone is broken up into nine belts (two belts for the crown and one each for the lands and grooves). Circumferentially, each belt is divided into ten segments. The result is determined as the arithmetic mean value within  $\pm 10\%$ .

The color Tsu of the deposits on the piston skirt is determined in the same way as the above index and is reproducible to within  $\pm 20\%$ . The piston skirt is divided into five belts.

In addition, the quality of the oil was evaluated on the basis of the following indices:

the amount of deposits  $K_0$  on the piston in g, reproducibility  $\pm 30\%$ ;

clogging of holes in oil-control rings and piston drainage holes  $Z_{mk}$  in points - from 0 (clean) to 10 (completely blocked), reproducibility  $\pm 30\%$ ;

wear index  $I_1$  in points, reproducibility  $\pm 30\%$ ;

wear of top ring  $C_{vk}$  in  $\mu$ , reproducibility  $\pm 40\%$ ;

corrosion index  $I_k$  in points, reproducibility  $\pm 10\%$ ;

the corrosive aggressiveness of the oil, determined visually and by weighing plates of type S-1 lead (k<sub>S-1</sub>), SOS 6-6 lead babbitt (k<sub>SOS</sub> 6-6), SB-30 lead bronze (k<sub>SB-30</sub>), and ASM aluminum alloy (k<sub>ASM</sub>), in  $\text{g}/\text{m}^2$ , reproducibility  $\pm 30\%$ ;

the stability index  $I_m$  of the oil in points, reproducibility  $\pm 20\%$ ;

the increase  $U_V$  in oil viscosity, determined as a percentage at  $50^\circ\text{C}$ , reproducibility  $\pm 15\%$ .

The indicators listed above are determined by the formulas

$$I_m = \frac{M_0 + M_1 + M_2 + M_3}{M_0 + M_1 + M_2 + M_3 + M_4 + M_5 + M_6 + M_7 + M_8 + M_9 + M_{10}} \cdot 100$$

$$M_1 = k_1 \cdot M_0 + k_2 \cdot M_1 + k_3 \cdot M_2 + k_4 \cdot M_3 + k_5 \cdot M_4 + k_6 \cdot M_5 + k_7 \cdot M_6 + k_8 \cdot M_7 + k_9 \cdot M_8 + k_{10} \cdot M_9$$

Table 2. Results of Tests of AS-6 011 with Control, Standard Method

a. Color marks	b. Deposits		c. Wear		d. Corrosion		e. Ring mobility		f. Stability index	
	L	S	1	2	3	4	5	6	7	8
b. Базисные	У	Прямоугольник	У	Прямоугольник	У	Прямоугольник	У	Прямоугольник	У	Прямоугольник
в. МА	Б	по окружности	Б	по окружности	Б	по окружности	Б	по окружности	Б	по окружности
г. МБ	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ
д. МВ	Б	С-Серия 1	Б	С-Серия 1	Б	С-Серия 1	Б	С-Серия 1	Б	С-Серия 1
е. МЕ	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ
ж. МА	Б	Призма	Б	Призма	Б	Призма	Б	Призма	Б	Призма
з. МБ	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ
и. МС	Б	ЦИАТИЧАСТЬ-1	Б	ЦИАТИЧАСТЬ-1	Б	ЦИАТИЧАСТЬ-1	Б	ЦИАТИЧАСТЬ-1	Б	ЦИАТИЧАСТЬ-1
к. МН	Б	ПМС-300	Б	ПМС-300	Б	ПМС-300	Б	ПМС-300	Б	ПМС-300
л. МС	Б	Серия 1	Б	Серия 1	Б	Серия 1	Б	Серия 1	Б	Серия 1
м. МС	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ	Б	СХ-Диагональ
н. МВ	Б	С-Серия 1	Б	С-Серия 1	Б	С-Серия 1	Б	С-Серия 1	Б	С-Серия 1

\*Plates of S-1, SOS 6-6, SB-30, and ASM.

- (a) Oil type
- (b) in USSR classification
- (c) in International classification
- (d) additive content, %
- (e) by weight
- (f) motor index, points
- (g) indicators included in motor index
- (h) deposits index, points
- (h<sub>1</sub>) indicators included in deposits index
- (i) ring mobility, points
- (j) color of deposits in ring zone, points
- (k) color of deposits on piston skirt, points
- (l) amount of deposits on piston, g
- (m) clogging of holes in oil control rings, points
- (n) wear index, points
- (o) spring weight, g
- (p) corrosion index, points
- (q) plate corrosion,  $\varepsilon/\text{h}^2$
- (r) stability index of oil, points
- (s) 50°C viscosity increase, %
- (t) remarks

and Recommended Additive Compositions on IKM-1 Machine by IKM-10FG

Additive composition number		h		t	
h	composition number	base	regular	premium	control
(1)	1	1	1	1	1
(2)	2	1	1	1	1
(3)	3	1	1	1	1
(4)	4	1	1	1	1
(5)	5	1	1	1	1
(6)	6	1	1	1	1
(7)	7	1	1	1	1
(8)	8	1	1	1	1
(9)	9	1	1	1	1
(10)	10	1	1	1	1
(11)	11	1	1	1	1
(12)	12	1	1	1	1
(13)	13	1	1	1	1
(14)	14	1	1	1	1
(15)	15	1	1	1	1
(16)	16	1	1	1	1
(17)	17	1	1	1	1
(18)	18	1	1	1	1
(19)	19	1	1	1	1
(20)	20	1	1	1	1
(21)	21	1	1	1	1
(22)	22	1	1	1	1
(23)	23	1	1	1	1
(24)	24	1	1	1	1
(25)	25	1	1	1	1
(26)	26	1	1	1	1
(27)	27	1	1	1	1
(28)	28	1	1	1	1
(29)	29	1	1	1	1
(30)	30	1	1	1	1
(31)	31	1	1	1	1
(32)	32	1	1	1	1
(33)	33	1	1	1	1
(34)	34	1	1	1	1
(35)	35	1	1	1	1
(36)	36	1	1	1	1
(37)	37	1	1	1	1
(38)	38	1	1	1	1
(39)	39	1	1	1	1
(40)	40	1	1	1	1
(41)	41	1	1	1	1
(42)	42	1	1	1	1
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(59)	59	1	1	1	1
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(61)	61	1	1	1	1
(62)	62	1	1	1	1
(63)	63	1	1	1	1
(64)	64	1	1	1	1
(65)	65	1	1	1	1
(66)	66	1	1	1	1
(67)	67	1	1	1	1
(68)	68	1	1	1	1
(69)	69	1	1	1	1
(70)	70	1	1	1	1
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(72)	72	1	1	1	1
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(74)	74	1	1	1	1
(75)	75	1	1	1	1
(76)	76	1	1	1	1
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(78)	78	1	1	1	1
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(80)	80	1	1	1	1
(81)	81	1	1	1	1
(82)	82	1	1	1	1
(83)	83	1	1	1	1
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(97)	97	1	1	1	1
(98)	98	1	1	1	1
(99)	99	1	1	1	1
(100)	100	1	1	1	1
(101)	101	1	1	1	1
(102)	102	1	1	1	1
(103)	103	1	1	1	1
(104)	104	1	1	1	1
(105)	105	1	1	1	1
(106)	106	1	1	1	1
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(108)	108	1	1	1	1
(109)	109	1	1	1	1
(110)	110	1	1	1	1
(111)	111	1	1	1	1
(112)	112	1	1	1	1
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(114)	114	1	1	1	1
(115)	115	1	1	1	1
(116)	116	1	1	1	1
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(128)	128	1	1	1	1
(129)	129	1	1	1	1
(130)	130	1	1	1	1
(131)	131	1	1	1	1
(132)	132	1	1	1	1
(133)	133	1	1	1	1
(134)	134	1	1	1	1
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(136)	136	1	1	1	1
(137)	137	1	1	1	1
(138)	138	1	1	1	1
(139)	139	1	1	1	1
(140)	140	1	1	1	1
(141)	141	1	1	1	1
(142)	142	1	1	1	1
(143)	143	1	1	1	1
(144)	144	1	1	1	1
(145)	145	1	1	1	1
(146)	146	1	1	1	1
(147)	147	1	1	1	1
(148)	148	1	1	1	1
(149)	149	1	1	1	1
(150)	150	1	1	1	1
(151)	151	1	1	1	1
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(165)	165	1	1	1	1
(166)	166	1	1	1	1
(167)	167	1	1	1	1
(168)	168	1	1	1	1
(169)	169	1	1	1	1
(170)	170	1	1	1	1
(171)	171	1	1	1	1
(172)	172	1	1	1	1
(173)	173	1	1	1	1
(174)	174	1	1	1	1
(175)	175	1	1	1	1
(176)	176	1	1	1	1
(177)	177	1	1	1	1
(178)	178	1	1	1	1
(179)	179	1	1	1	1
(180)	180	1	1	1	1
(181)	181	1	1	1	1
(182)	182	1	1	1	1
(183)	183	1	1	1	1
(184)	184	1	1	1	1
(185)	185	1	1	1	1
(186)	186	1	1	1	1
(187)	187	1	1	1	1
(188)	188	1	1	1	1
(189)	189	1	1	1	1
(190)	190	1	1	1	1
(191)	191	1	1	1	1
(192)	192	1	1	1	1
(193)	193	1	1	1	1
(194)	194	1	1	1	1
(195)	195	1	1	1	1
(196)	196	1	1	1	1
(197)	197	1	1	1	1
(198)	198	1	1	1	1
(199)	199	1	1	1	1
(200)	200	1	1	1	1
(201)	201	1	1	1	1
(202)	202	1	1	1	1
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(206)	206	1	1	1	1
(207)	207	1	1	1	1
(208)	208	1	1	1	1
(209)	209	1	1	1	1
(210)	210	1	1	1	1
(211)	211	1	1	1	1
(212)	212	1	1	1	1
(213)	213	1	1	1	1
(214)	214	1	1	1	1
(215)	215	1	1	1	1
(216)	216	1	1	1	1
(217)	217	1	1	1	1
(218)	218	1	1	1	1
(219)	219	1	1	1	1
(220)	220	1	1	1	1
(221)	221	1	1	1	1
(222)	222	1	1	1	1
(223)	223	1	1	1	1
(224)	224	1	1	1	1
(225)	225	1	1	1	1
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(228)	228	1	1	1	1
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(230)	230	1	1	1	1
(231)	231	1	1	1	1
(232)	232	1	1	1	1
(233)	233	1	1	1	1
(234)	234	1	1	1	1
(235)	235	1	1	1	1
(236)	236	1	1	1	1
(237)	237	1	1	1	1
(238)	238	1	1	1	1
(239)	239	1	1	1	1
(240)	240	1	1	1	1
(241)	241	1	1	1	1
(242)	242	1	1	1	1
(243)	243	1	1	1	1
(244)	244	1	1	1	1
(245)	245	1	1	1	1
(246)	246	1	1	1	1
(247)	247	1	1	1	1
(248)	248	1	1	1	1
(249)	249	1	1	1	1
(250)	250	1	1	1	1
(251)	251	1	1	1	1
(252)	252	1	1	1	1
(253)	253	1	1	1	1
(254)	254	1	1	1	1
(255)	255	1	1	1	1
(256)</					

Table 3. Results\* of Tests of Soviet Oils from Eastern Petroleum

No.	Name	b) $T_{40}$ (°C)				c) Motor index				d) Deposits				e) Viscosity				
		AS-6	AS-6 + 0.5% PMS-200A	AS-6 + 1% PMS-200A	AS-6 + 1.5% PMS-200A	AS-6	AS-6 + 0.5% PMS-200A	AS-6 + 1% PMS-200A	AS-6 + 1.5% PMS-200A	AS-6	AS-6 + 0.5% PMS-200A	AS-6 + 1% PMS-200A	AS-6 + 1.5% PMS-200A	AS-6	AS-6 + 0.5% PMS-200A	AS-6 + 1% PMS-200A	AS-6 + 1.5% PMS-200A	
1	Opetna, Gasoline	6.8	7.2	9.6	9.8	9.8	9.8	9.8	9.8	6.8	7.2	9.6	9.8	6.8	7.2	9.6	9.8	
2	Motor oil, regular	3.8	4.0	9.0	9.2	9.5	9.8	9.8	9.8	3.8	4.0	9.0	9.2	3.8	4.0	9.0	9.2	
3	Motor oil, premium	4.2	4.5	4.8	6	3	13	4	4.2	4.5	4.8	6	3	13	4	4.2	4.5	4.8
4	Motor oil, heavy-duty	4.2	4.5	4.8	6	3	13	4	4.2	4.5	4.8	6	3	13	4	4.2	4.5	4.8
5	Motor oil, MB	4.2	4.5	4.8	6	3	13	4	4.2	4.5	4.8	6	3	13	4	4.2	4.5	4.8
6	Motor oil, MV	4.2	4.5	4.8	6	3	13	4	4.2	4.5	4.8	6	3	13	4	4.2	4.5	4.8
7	Santolube 493	57.5	57.5	47.5	40.1	47.1	46.4	46.4	57.5	57.5	47.5	40.1	47.1	46.4	57.5	57.5	47.5	40.1
8	Santolube 493; (g) rating, points	67.7	67.7	57.5	47.5	40.1	47.1	46.4	67.7	67.7	57.5	47.5	40.1	47.1	67.7	67.7	57.5	47.5
9	Piston crown	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—
10	Viscosity, mg	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—
11	Viscosity increase at 37.8°C, %	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—	—

\*Obtained by Monsanto by JP 175/60T method on Pittier W-1 engine.

KEY: (a) indicator; (b) group (series) of oil and additive composition; (c) base (regular); (d) specimen... (e) MA (premium) + 0.7% Monto 613 + 0.7% Santolube 493; (f) MB (Heavy Duty) + 1.5% Monto 613 + 0.7% Santolube 493; (g) MV (series 1) + 4% Monto 613 + 0.25% Santolube 493; (h) rating, points; (i) piston skirt; (j) piston crown; (k) mass loss of lead-bronze bearing, mg; (l) viscosity increase at 37.8°C, %.

The color of the deposits in the piston-ring zone was 6.9 points on the VNI NF scale in a test of AS-6 oil with 0.5% of TSATIM-339 additive and 0.5% PMS-Ya, but when 0.005% of PMS-200A was added to the same mix, it was lowered to 4.6 points. The same result was obtained on introduction of 0.005% of PMS-200A into an additive mix consisting of 3.5% VNI NP-360 and 1% AZNII-TSATIM-1 in AS-6 oil. In this case, the color of the deposits in the piston-ring zone is lowered from 8 to 5.8 points. This makes it possible to bring GOST 10541-63 Group MB oils up to the level of group MV, improves piston-ring conditions, and reduces piston deposits.

The thermal stability of oil with these additive mixes was also improved, as is clearly indicated by the decrease in the rate of viscosity change in tests of the oil on the IIM-1 machine. For example, while the additive mix consisting of 1% TSATIM-339 and 0.5% PMS-Ya caused a 13.9% viscosity increase in AS-6 oil, this increase was only 5.6% when 0.005% of PMS-200A was added.

On introduction of the additive mix consisting of 3.5% VNI NP-360 and 1% AZNII-TSATIM-1 into AS-6 oil, viscosity increases by 18.1%; with simultaneous addition of PMS-200A, the viscosity increases by only 5.4%.

This improvement of the use properties of the oil on introduction of the chain-stopping additive into a mix containing

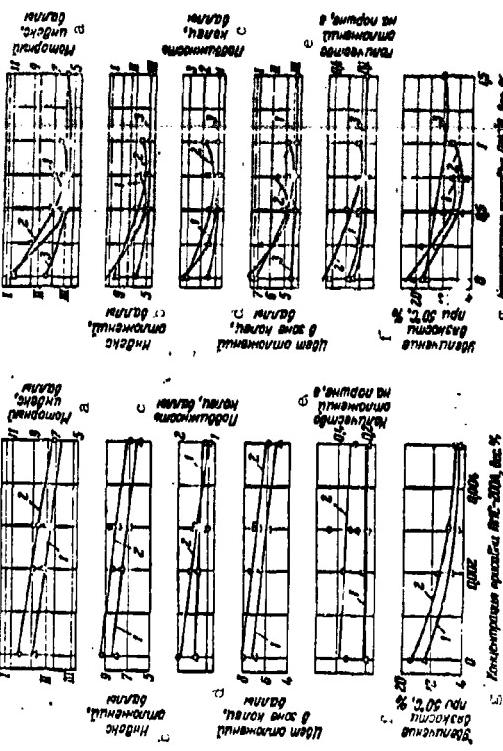


Fig. 2. Influence of content of PMS-200A additive on motor properties of AS-6 oil with the following additive compositions: 1) AS-6 + 1% TSATIM-339 + PMS-Ya + 0.005% PMS-200A; 2) AS-6 + 1.5% TSATIM-339 + PMS-Ya + 0.05% PMS-200A; 3) AS-6 + 2.5% TSATIM-339 + PMS-Ya + 0.5% PMS-200A. I) Level of group MA control oil; II) same, group MB; III) same, group MV.  
KEY: (a) motor index, points; (b) deposits index, points; (c) ring mobility, point; (d) color of deposits in ring zone, points; (e) quantity of deposits on piston, %; (f) increase in viscosity at 50°C, %; (g) concentration of PMS-Ya additive, % weight.

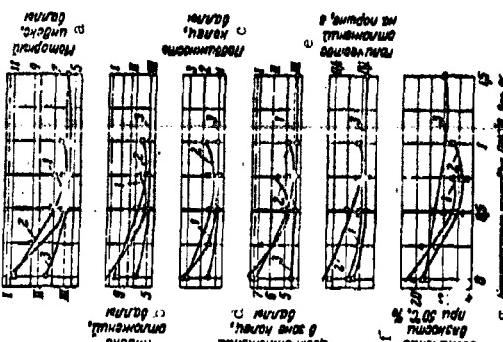


Fig. 3. Influence of content of PMS-Ya additive on motor properties of AS-6 oil with the following additive compositions: 1) AS-6 + 1% TSATIM-339 + PMS-Ya + 0.005% PMS-200A; 2) AS-6 + 1.5% TSATIM-339 + PMS-Ya + 0.05% PMS-200A; 3) AS-6 + 2.5% TSATIM-339 + PMS-Ya + 0.5% PMS-200A. I) Level of group MA control oil; II) same, group MB; III) same, group MV.  
KEY: (a) motor index, points; (b) deposits index, points; (c) ring mobility, point; (d) color of deposits in ring zone, points; (e) quantity of deposits on piston, %; (f) increase in viscosity at 50°C, %; (g) concentration of PMS-Ya additive, % weight.

detergent and antioxidant components can be explained by the decreased aeration of the oil, which has a strong influence on quality changes of oil in the engine.

A test of TsLATIM-339 additive in concentrations of 1.0%, 1.5, and 2.5% in a mix with 0.005% PMS-200A indicated (Fig. 3) that the amount of piston deposits decreases from 0.65 to 0.32 g, ring mobility from 1.5 to 1 point, and the color of the deposits in the piston-ring zone from 7.2 to 4.5 points as the concentration of the TsLATIM-339 additive is increased to 2.5%. In this case, the motor index drops from 11 points (the level of the group MA control oil) to 7.5 points, which corresponds to the level of MB control oil. As the test results showed, the tested concentrations of TsLATIM-339 additive do not provide adequate detergent properties in the oil, and for this reason they were supplemented with PMS'Ya detergent-dispersing additive in concentrations of 0.25, 0.5, 0.75, 1, 1.25, and 1.5% by weight.

On introduction of small amounts of the new additive mixes, ring mobility drops to 1 point, the color of deposits in the piston-ring zone to 4.2-4.5 points, the amount of piston deposits to 0.16-0.2 g, and the rate of oil viscosity increase after the test to 5-8%. In this case, the motor index declines to 5.5-6 points, which corresponds to the level of the Group MV control oil (series 1). However, it must be noted that the optimum values can be obtained only at definite proportions of the TsLATIM-339 and PMS'Ya additives, in the 2:1 range. This ratio also applies for diesels operated on low-sulfur fuels (Fig. 4). Departure from it results in a rise of the motor index, indicating deteriorating oil quality.

Adjustment of the mix on the basis of the TsLATIM-339, PMS'Ya, polymethacrylate D (for winter-grade oils), and PMS-200A additives, which was carried out on the IKM-1 machine by the IKM-10FG method, resulted in tentative recommendations of oils conforming to the Premium, Heavy-Duty, and Series 1 groups. The results of tests on these oils were compared with those for oils with the imported control additives Monto 613 and Santolube 493.

The above additive mixes were also tested on a machine with a full-scale Moskvich 407 automobile engine, and the correctness of the results obtained was confirmed.

Thus, the additive mix given in Table 4 can be recommended on the basis of the tests for carburetor-type automobile engines.

Tests of another additive mix, consisting of ASK, DF-11, polymethacrylate D (for winter oils) and PMS-200A in AS-6 oil, were run by the IKM-10FG method on the IKM-1 installation. The content of the ASK additive in the mix was varied from 1.5 to 3.5%. As the tests showed (Fig. 5), as little as 1.5% of ASK produces an oil conforming to Group MB (Heavy Duty); when the ASK concentration is increased to 2.5%, Group MV (Series 1) oils

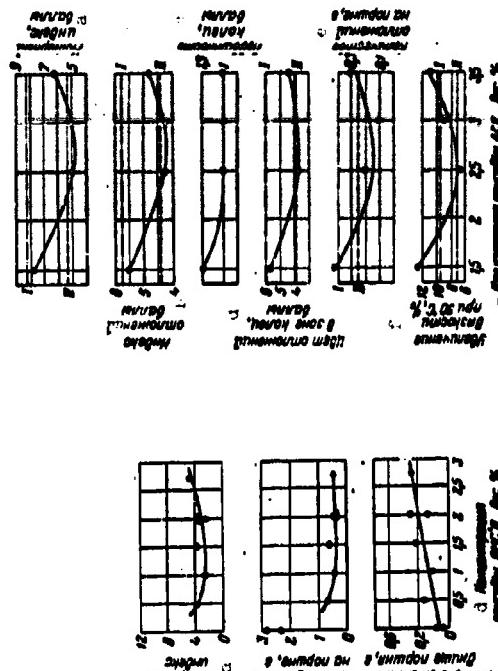


FIG. 4. Influence of content of PMS'Ya additive on motor properties of oil with additive mix consisting of 3% TsLATIM-339, PMS'Ya, and 0.005% PMS-200A.  
KEY: (a) motor index; (b) weight of deposits on piston; (c) weight of head deposits on piston head; (d) PMS'Ya additive concentration, % by weight.

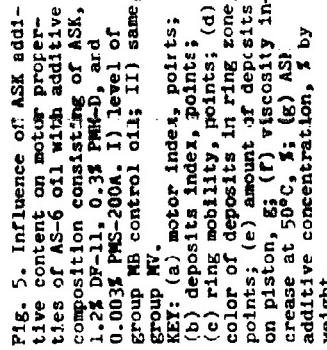


FIG. 5. Influence of ASK additive content on motor properties of AS-6 oil with additive composition consisting of ASK, DF-11, 0.3% PMS-D, and 1.2% PMS-200A.  
KEY: (a) motor index, points; (b) weight of deposits on piston, g; (c) ring mobility, points; (d) color of deposits in ring zone, points; (e) amount of deposits on piston, g; (f) viscosity increase at 50°C, %; (g) ASK additive concentration, % by weight.

are obtained. Here the color of the piston deposits drops from 5.7 to 3.7 points on the VNII NP scale, the amount of deposits on the piston is reduced from 0.43 to 0.1-0.2 g, and the degree of viscosity increase also drops from 12.9 to 6.4%. The decrease in the motor index from 7.8 to 5 points also indicates an improvement of the operational properties of oil with these additives.

Table 4. Recovery of  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$

Примечание	Состав гидролизата, мг/л		
	CH <sub>3</sub>	SH	NH
ИУАТИМ-329	0,1-1	4-15	1,5-2
ИУАТИМ-329	0,25-0,5	0,75-1	1-1,5
Индустритовая А (Хим. завод Азота)	0,3	3%	0,3
ИУАТИМ-329	0,1-0,5	0,1-0,5	0,025
ИУАТИМ-329	0,1-0,5	0,1-0,5	0,025

Figure 1(a) illustrates the effect of the number of hidden neurons on the error.

Table 5. Parameters of model 3.

Применение	Задание на проверку	
	Проверка	Норма
АСК АФ-11 на ВИНИ НИК-35	4-15 0,6-1,2	1,5-2,5 0,6-1,2
Пневмогазорез Д (для замены ма- териала)	0,3 0,003	0,3 0,003
ПМС-200A		

**KEY:** (a) additive; (b) additive component; (c) by-weight; (d) by-volume; (e) by-weight (f) ASK; (g) or until further notice; (h) permanent; (i) winter-grade oil; (j) -in-GA.

In vacuums the concentration of  $\text{AgP}$  additive to  $\text{Si}_3\text{N}_4$  in this glass increases the thermal stability of the glass and the  $\text{AgP}$  index rises from 4.5 to 5.5 points.

Oil with these additive mixes were bench-tested in a multi-cylinder Mackin 407 automobile engine. The test results showed

The test results are a basis for recommending one more adhesive mix (1% by weight) for oils used in carburetor-type automobile engines (Table 5).

SNC SII 203

1. A method designated IEM-10FG was developed for motor rating of the base properties of base oils and additive oils of Groups IIA (Pump Oil), ME (Heavy Duty), and MV (Ferries) on a special test rig.

## References

其後又復有大風雨，人謂之爲「天子氣」。

THE CHAMBERS OF COMMERCE - THE USE OF PRO-  
THE CHAMBERS OF COMMERCE - THE USE OF PRO-

Scientific Lubrication, Cleveland, Ohio, The U.S. Steel Co., Pittsburgh, Pa., and Standard Oil Co., New York, N.Y.

Ball, J.A., Scherren, S.M. and Nelson, J.B. New test technique for aircraft piston engines, SAE Trans., No. 1263.

## References

- Notice  
of a meeting of the Board of Directors of the First National Bank of Marion, Ohio, to be held at the office of the bank, Marion, Ohio, on Friday, April 13, 1917, at 10 o'clock A.M.

Symbol List

## TESTS OF SOVIET AND FOREIGN MOTOR OILS IN THE UIM-6-NAII AND UIM-6N-NATTI SINGLE-CYLINDER BENCH INSTALLATIONS

S.G. Arabyan, G.P. Belyaevskiy, I.N. Danilov, V.V. Makanarov,  
I.A. Kholemonov, O.N. Sneyrina, Ye.N. Pirsarova

The object of the study reported in this paper was to design a single-cylinder tractor built on a series-produced supercharged tractor diesel, develop a test method, and rate various types of motor oils.

The properties of oils intended for work under relatively light-duty conditions (Premium type) have been rated on the UIM-1 single-cylinder torch built at the Autonole and automotive Engine Scientific Research Institute (NATI) or the basis of the F-5 tractor diesel. Later, the UIM-6 NATI machine was developed on the basis of the D-75 tractor diesel, which is a supercharged modification of the D-54, for testing Heavy Duty and Series 1

For rating Series 2 cells, the apparatus was fitted with a surecharging unit that delivered a substantial increase in horsepower per liter; this installation was designated UIM-6N-MAP (see figure).

Below we list the technical details of our 6N-MATI installations.

UIM-6N-MATI	
Power, hp.....	73
Granthaft speed, rev/min.....	1530
Net effective structure, kg/cm <sup>2</sup> .....	9.7
Horsepower per liter.....	16.1
Compression ratio.....	16
Bore, mm.....	125
Stroke, mm.....	152
Max. cylinder pressure, kg/cm <sup>2</sup> .....	0.5
Fuel consumption, kg/hour.....	5
Oil injection system capacity, kg.....	5
Specific lubricating system capacity, kg/hap.....	0.17
Concentrated heating index of cylinder-firing group.....	71
Concentrated oil-duty index.....	426
Lubricating system.....	21
Cooling system.....	20
Coilants.....	19

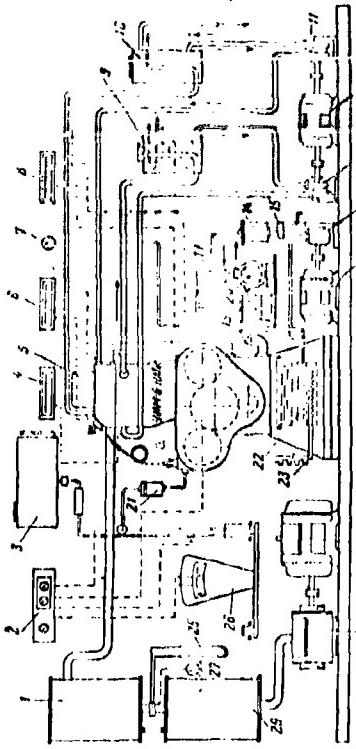


FIG. 1. Diagram of UIM-6N-MATI bench installation for PSID-ZM testing. 1, 19, 29) receivers; 2) instrument for measuring fuel flow rate; 3) PSID-ZM-MATI; 4, 6, 8) lagometers [sic]; 5) engine; 9) heat exchanger; 10) mixing tank; 11, 13) water pumps; 12, 18, 24) electric motors; 14) oil filter; 15) reduction valve; 16) oil pump; 17) oil radiator; 20) gas meter; 21) fuel filter; 22) oil sump; 23) LATR [lab autotransformer]; 25) piezometer; 26) scales; 27) thermistor; 28) supercharger. KEY: (a) UIM-6-MATI.

In developing the method for rating the motor properties of crankcase oils from their tendency to form carbon deposits, we chose engine operating conditions under which the type and degree of parts fouling by deposits formed in full-scale engines during long-term use could be duplicated during a relatively short test time.

#### TEMPERATURE MEASUREMENTS ON THE ENGINE

The temperature condition of the components of the cylinder-piston group has a strong influence on the process of deposit formation in an engine; for this reason, temperature readings were taken at the sleeve and piston of the UIM-6-MATI installation under various engine operating conditions.

It was established that the highest piston-head temperature (at the center of the combustion chamber) in operation without supercharging and at an effective pressure of  $P_e = 6.8 \text{ kg/cm}^2$  is  $700^\circ\text{C}$ , while that on the side of the piston head is  $200^\circ\text{C}$  lower. The temperatures in the areas of the first and second grooves are  $600^\circ\text{C}$  and  $520^\circ\text{C}$ , respectively. Evidently, caused no substantial

change in the temperature distribution on the piston and cylinder sleeve; at  $P_e = 9.5 \text{ kg/cm}^2$ , the temperature rises by only  $10-15^\circ\text{C}$ .

Similar results, which are presented on page 65, were obtained on the SMD-14, AM-01, SMD-18, and YAMZ-238MV [sic] engines. When glycerine was used as a coolant instead of water, the temperatures of the engine parts rise sharply. For the first and second piston grooves, this temperature increase is  $550^\circ\text{C}$ .

Piston temperature increases by  $0.5^\circ\text{C}$  on a  $1^\circ\text{C}$  increase in the glycerine temperature.

#### INFLUENCE OF ENGINE TEMPERATURE CONDITIONS ON RATE OF CARBON-SCALE FORMATION

It has been established [1] that the rate of fouling of engine parts is determined in many respects by engine-temperature conditions and test time; the test time can be shortened by increasing the heat loss on the engine part. However,

Table 1. Influence of Coolant Temperature on Rate of Piston Fouling

Series	Type of piston	Mean effective pressure, atm.	Average water-coolant ratios, $\frac{\text{kg}}{\text{liter}}$			Water temperature, °C	Rate of piston fouling, points
			10	20	30		
1	Cast iron piston with two grooves	1.30	1.60	1.80	1.90	135	—
2	Glycerine piston with two grooves, $\text{kg}/\text{liter}$	6.5	6.3	5.6	5.7	75	—
3	Glycerine piston with two grooves, $\text{kg}/\text{liter}$	11.3	9.5	9.4	15	45	—
4	Cast iron piston with two grooves, $\text{kg}/\text{liter}$	—	—	—	0.5	0.35	—
5	Piston rings	0.25	0.22	0.17	0.21	0.13	—
6	Cast iron piston with two grooves, $\text{kg}/\text{liter}$	—	—	—	—	—	—
7	Glycerine piston with two grooves, $\text{kg}/\text{liter}$	—	—	—	—	—	—
8	Cast iron piston with two grooves, $\text{kg}/\text{liter}$	34.2	49.2	40.5	74	69.5	55.6
9	Cast iron piston with two grooves, $\text{kg}/\text{liter}$	212	192	192	435	292	63
10	Polypropylene piston, $\text{kg}/\text{liter}$	—	—	—	—	—	—

\*By UIM-6-NATI method.

\*\*Rated in points as a function of the degree of mobility of each piston ring. Here the condition of a ring that can be moved freely in its groove is given a point rating of 0; the condition of a ring that is coked around its entire circumference is given a rating of 10.

\*\*\*Rated in points as a function of the amount and type of deposits (semifluid, hard, etc.) in each groove. A perfectly clean groove is given a rating of 0; a groove completely filled with solid deposits is rated 10.

\*\*\*\*The point ratings characterizing the fouling of all piston zones are totaled. In this table and in Tables 2, 3, and 6, the indicators (in points) characterize the presence of deposits on the piston skirt, on the inside surface of the piston, etc., are not listed, although they are included in the over-all rating.

KEY: (a) UIM-6-NATI; (b) SNC-1-45; (c) UIM-6-NATI; (d) VIM-6-NATI; (e) horsepower; (f) crankshaft speed, rev/min.; (g) mean effective pressure, kg/cm<sup>2</sup>; (h) sp./liter; (i) total pressure, kg/cm<sup>2</sup>; (j) specific lubrication system viscosity, kg/cm<sup>2</sup> sec.; (k) conventional oil index of cylinder-piston temperature; (l) piston temperature; (m) first groove; (n) second groove.

slight increase in engine thermal conditions may change the direction of the piston rings, with the result that objective rating of the oil's quality is distorted.

Therefore, taking a 120-hour test time, which corresponds to the average oil-change interval in tractor engines, we tested heavy-duty oil at a water-coolant temperature of 96°C and glycerine temperatures of 96, 115, and 135°C to find the optimum thermal regimes on the UIM-6-NATI bench. The test results in Table 1 indicate that scaling on the piston increases substantially as running temperatures rise, especially when we pass to the 135°C glycerine temperature, and the over-all piston-fouling rating reaches 16.2 points.

With the glycerine temperature rises to 135°C, the rate of deposit accumulation decreases and the over-all rating rises only 3 points.

KEY: (a) Indicators; (b) Heavy Duty (UIM-6-NATI); (c) water; (d) Glycerine; (e) Series 1 (UIM-6-NATI); (f) Series 2 (UIM-6-NATI); (g) cast iron piston; (h) cast iron piston-skirt sealing in piston-groove, points; (i) over-all carbon scaling rating for piston, points; (j) amount of deposits in piston-groove and on piston, g.

Similar results were obtained in tests of Series 1 and Series 2 oils on the UIM-6-NATI bench at various coolant temperatures. In the tests of Series 2 oil, going from 115 to 135°C is detrimental to the mobility of the first compression ring and reduces piston fouling slightly, probably because of burning-out of the deposits in the ring grooves.

On the basis of the results obtained on piston fouling and the physicochemical properties of the used oils, the coolant (glycerine) temperature was set at 115°C. At this temperature, Heavy Duty and Series 1 oils are differentiated quite sharply on the UIM-6-NATI machine (16.2 and 9.8 points, respectively); at the same temperature, the over-all piston-scaling figures obtained on the UIM-6-NATI bench for Series 1 and Series 2 oils differ even more widely (20.5 and 12 points, respectively); also, the first compression ring was coked after 60-80 hours of the test on Series 1 oil.

Thus, the test conditions arrived at made it possible to classify Heavy Duty, Series 1, and Series 2 oils clearly and reliably.

#### REPRODUCIBILITY OF TEST RESULTS

One of the important stages in developing an oil-test method is to check the reproducibility of the results of parallel experiments.

Table 2. Results of Parallel Tests on Heavy Duty and Series 1 Control Oils

a Indicators (in points)	b Xenon-Toluene		c Cetene	
	0.2	4.8	expansion coefficient, %	expansion coefficient, %
e Nitrope magnesia nozoch f Harpoorotene magnesia nozoch g Harpoorotene magnesia nozoch h Cetene oxygen magnesia nozoch	0 14 11 4.8	0 10 0.3 6.6	0 9.2 29 16.2	0 4.6 2.1 9.8
i j k l m n o p q r s t u v w x y z	0 14 11 4.8	0 10 0.3 6.6	0 9.2 29 16.2	0 4.6 2.1 9.8

KEY: (a) indicator (in points); (b) Heavy Duty; (c) Series 1; (d) sulfur content in fuel; (e) piston ring mobility loss; (f) scaling in first ring groove; (g) scaling in other grooves; (h) over-all piston-scale rating.

For this purpose, repeated tests were made with Heavy Duty, Series 1, and Series 2 oils both on the same installations and on several installations at the N.A.T.I., the All-Union Scientific Research Institute of Oil and Gas Processing and Production of Artificial Liquid Fuel (VNI NP), the Eastikir Scientific Research Institute of Petroleum Refining (BashNII NP), and the Institute of Petrochemical Processes (INKh AN) of the Azerbaijan SSR.

The test results, which appear in Tables 2 and 3, indicate satisfactory agreement between the results of the tests on all oil types.

Table 3. Results of Parallel Tests of Series 2 Control Oil

a Indicators (in points)	b Xenon		c Cetene		d Sulfur content in fuel		e Harpoorotene magnesia nozoch		f Harpoorotene magnesia nozoch		g Cetene oxygen magnesia nozoch		h Cetene oxygen magnesia nozoch	
	r	n	i	w	t	u	v	x	y	z	w	u	v	x
f Nitrope magnesia nozoch g Harpoorotene magnesia nozoch h Cetene oxygen magnesia nozoch	0 14 11 4.8	0 10 0.3 6.6	0 9.2 29 16.2	0 4.6 2.1 9.8	0 1.5 3.0 1.6	0 3.5 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5
i j k l m n o p q r s t u v w x y z	0 14 11 4.8	0 10 0.3 6.6	0 9.2 29 16.2	0 4.6 2.1 9.8	0 1.5 3.0 1.6	0 3.5 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5	0 3.4 0.7 0.5

KEY: (a) indicators (in points); (b) BashNII NP; (c) VNI NP; (d) INKh AN; (e) INKh NP; (f) loss c.; (g) scaling in first ring groove; (h) scaling in other grooves; (i) over-all piston-scale rating.

COMPARISON OF OIL-TEST RESULTS OBTAINED ON SINGLE-CYLINDER BENZINE AND ON TRACTOR DIESELS

In many cases, the results of oil tests on certain laboratory and motor machines do not agree with results obtained on full-sized production engines. This is chiefly because the operating and test conditions set up on the laboratory and motor installations do not match the design features and operating conditions of the production engines.

To one degree or another, almost all of the methods that have been developed permit classification of oils, but many of them have not been used extensively because tests of the same oils in engines give inconsistent results. Hence the basic criteria for establishing the reliability of the results should be the results obtained in long-term tests of the same oil specimens on the most common types of engines.

Tests were run on Heavy Duty (with low-sulfur and sulfur-containing fuels) and Series 1 control oils on the UIM-6-NATI bench and in SMD-14 and D-28 tractor diesels in order to determine the reproducibility of the results.

The indicators listed in Table 4 indicate that the high-temperature test regime selected for the UIM-6-NATI bench, while it made it possible to reduce the test time to 120 hours, enables one to obtain the same results as with production engines operating under normal temperature conditions for 500 hours. The UIM-6-NATI installation occupies an intermediate position between the SMD-14 and D-28 engines as regards almost all indicators of piston and ring fouling by carbon deposits.



Glycerine in cylinder jacket 115  
oil in crankcase 95

We see from the results obtained (table 6) that D-311 with 5% VIII MP-360 has properties similar to those of the Heavy Duty control oil.

DS-11 oil with 3% TSITIM-315 and oil with 3% TSITIM-319 + 2% AFB have motor properties considerably below those of the Heavy Duty control oil. The same oil with the following additive mixes:

These use properties approaching those of the Series I control oil, while BS-II oil with an additive mix consisting of 3% ISIMIX 209 + 2% PMSV and 0.05% PMS-200A has anti-scaling properties slightly better than those of the control.

The test results obtained in the single-cylinder bench unit were in full agreement with the previous long-term tests run on multi-hole D-20 and GM-4 engines.

Table 7 lists the proportions of these oil-soluble additives to those of the Jensen conventional oil. A note should be made concerning the use of the high-use property of PMS Ya + 0.005% PMS-2004, which has given an overall rating of 14.5 compared to 13.0 for the control oil.

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1. A general-purpose single-cylinder bench engine designated URM-6-NAT and a supercharged modification, URM-6-SHAT, were tested and built for evaluation of the use properties of various types of oils made for use in existing and projected tractor diesels.
  2. A battery of tests was run to adjust test conditions and develop a technique for quick one-hour tests of heavy duty Series 1, and Series 2 motor oils.
  3. The tests showed that the conditions selected permit sharp differentiation of the oils and produce satisfactory

Table 5. Results of Tests of Oils with Domestic Additives

on UIM-6-NATI Bench and D-50 and SMD-14 Engines

Table 7. Results of Testing Series 2 Oils with Domestic Additives on UIM-6-NATI Machine

E	E	UIM-6-NATI Carburetor 1921 + Novosibirsk OILS AND FATS ZAO (C-pow 1)	UIM-6-NATI Carburetor 1921 + Novosibirsk OILS AND FATS ZAO (C-pow 1)											
			M	N	O	P	Q	R	S	T	U	V	W	X
UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI	UIM-6-NATI
0	1.25	0	0	0	0	0	0	0	0	0	0	0	0	0
10	6.5	5.5	5.2	2.7	4.5	4.5	3.1	2.8	6	6.4	1.7	6.3	—	—
20	6.8	2.5	4.9	2.5	2.9	1.5	0.9	2.6	2.3	3	2.9	—	—	—
212	6.8	20.9	10.5	9.2	10.8	12	10.5	11	12	16.9	5.2	10.3	—	—
2.9	4.6	1.1	1.3	1	0.9	1.2	1.6	0.9	1.4	1.6	2.5	1.04	—	—
22	0.5	3.1	0.8	0.6	1.1	0.9	1.15	0.9	1.25	1.4	0.65	1.14	—	—
6.5	2.1	5.6	2.6	3.1	3.9	4.3	2.1	2.7	3.2	3.4	1.2	0.5	—	—
0.5	1.3	0.5	1.1	1.35	0.4	1.1	1.9	0.7	1.2	0.7	1.2	0.5	—	—

(1) 3% Rashnyi-2 + 1% PMS Ya + (r) carbon scaling in other grooves; points;  
 (m) UIM-6-NATI  
 (n) SMD-14  
 (o) —  
 (p) piston-ring mobility loss,  
 (q) piston scaling in first ring, points;  
 (r) carbon scaling in first ring, points;

(s) piston wear, points;  
 (t) first-ring wear, points;  
 (u) wear of set of rings, i.e., groove, points;  
 (v) amount of scale in grooves and on rings, %;  
 (w) sleeve wear, i.e., wear of set of rings, E.

agreement and reproducibility between parallel experiments either on a single bench unit or a number of such units.

4. Test results obtained for various foreign and Soviet oil on the UIM-6-NATI and UIM-6-NATI machines agree with data obtained in long-term (500-800-hour) tests of the same oil samples in the D-50, SMD-14, D-28, SMD-18 and YMZ-238ND traction diesel.

5. Production and test samples of Soviet motor oils with additive compositions developed at the UIM NP, INKKP AN Azer. SSR and BashNII NP were classified.

6. The quick test methods developed for Heavy Duty and Service 1 oils were standardized, and devised for Service 1 to recommended for standardization and general use.

#### References

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#### OIL AND OIL-ADITIVE SELECTION AND USE-PROPERTY MOTOR RATING FOR TWO-STROKE GASOLINE ENGINES

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As we know, the oil that lubricates two-stroke gasoline engines is introduced through the carburetor together with the fuel.

Depending on the design, physical condition, and operating conditions of the engines and on the properties of the oil and fuel air, the oil-to-fuel ratio varies from 1:15 to 1:40.

Oil for lubrication of two-stroke gasoline engines must meet the following main requirements:

- 1) Not burn with formation of the smallest possible amount of carbon, while ensuring minimal wear of engine parts, longer oil engine operating conditions;
- 2) Not contain particles capable of plugging the engine fuel-supply system;
- 3) Must ensure normal lubrication of the engine at minimum oil-air ratios;
- 4) Must protect the parts of the engine and its fuel-supply system from corrosion;

5) Have good properties during storage is required when oil is mixed with additives and stabilizers;

In the event that the gasoline becomes watered, the oil and additives may not settle out of the mixture.

Oils with varying viscosities, chemical compositions, and additive contents are used in various countries to lubricate two-stroke gasoline engines.

It has been established that light type A-6 oils with medium viscosity indices produce the smallest amounts of deposits on the piston, in the combustion chamber, in the scavenging and exhaust port, and on spark-plug electrodes. While a heavier oil of type MS-19 with high viscosity index eliminates pitting and scoring of the pistons.

Use of mixed light and heavy oils that have been matched with consideration of chemical composition, the specific use conditions, and the engine design gives good results.

The use properties of the oils are improved considerably by introduction of special additives. Use of commercial and tractor oils with ashing additive is not recommended, since the additives increase wear and the amount of deposits in the scavenging and exhaust ports, in the combustion chamber, and on the plug electrodes.

It should be noted that factory recommendations on oils to be used in two-stroke gasoline engines are frequently given without adequate technical justification.

At the All-Union Scientific Research Institute of Oil and Gas Processing and Production of Artificial Liquid Fuel (VNII NF), development of special additive oils was begun with elaboration of a spot-testing method using the SD-60E engine. Here it was necessary that:

the test method require little time and no more than 0.5 kg of the experimental oil; and that, provide reproducible experimental results corresponding to the results of long-term bench and field tests of oils in two-stroke gasoline engines of various ratings.

The engine permit heat-loaded simulation of existing and projected two-stroke gasoline engines and provide sharp differentiation of oils on the basis of the main use indicators; further, that the engine be simple to operate and that it perform smoothly and reliably under various thermal loads, with adequate service life of main replacement of the basic components - the piston and cylinder.

Among the two-stroke gasoline engines that we have at the present time, the SD-60E meets these specifications, and was chosen as a basis for development of the bench installation and the spot testing method.

\*See Part 4 for footnote.

Unlike motor-vehicle and traction engines, it does not require an outside blower when operated under bench conditions. This adequate cooling fan makes it possible to vary the thermal stressing of the engine over a broad range by throttling the air flow and, if necessary, to simulate use conditions.

Below we give the technical data of the SD-60E engine.

Available power, hp.....	4000
Crankshaft speed, rev/min.....	air
Cooling.....	A-66 or B-70
Fuel.....	gasoline mixed with oil (25:1)
Fuel consumption, g/hp-h,	
not above.....	550
Weight of engine (without muffler), kg.....	7
Guaranteed service life, h.....	600
Weather climatic, altitude and amortization losses.	

In designing the oil-testing installation, certain auxiliary equipment was provided to allow for changes in application of the engine; they make it possible to control and monitor its operating conditions.

Essentially, the method consisted in a short test of the oil in the engine at high temperature. The motor qualities of the oils were rated on the basis of the amount of deposits on a piston, in the cylinder head, and in the cylinder exhaust port, and from the degree of piston-ring mobility.

The use properties of the oil were determined by comparing the results of tests run on the same installation with the test oil and a control oil, in terms of the rating indicators adopted. The control oils were base oils made according to the appropriate GOST [All-Union State Standard] and addition oils for two-stroke gasoline engines, which were prepared in accordance with the appropriate Technical Specifications [TSS] or GOSTs.

The control oil was tested after every 4-5 tests, run with experimental samples. This made it possible to check the technical condition of the bench, its capability for differentiation, and the identity of the experimental conditions.

The following test conditions were adopted as a result of the studies (320 E of oil are required per 1 experiment):

Test time, h	10
Crankshaft Speed, rev/min	4000±10
Compression ratio	7.1
Geometrical actual	5.3

Spark timing before top dead center, deg..... 27-28  
 Fuel... B-70 gasoline, GOST 1012-54  
 Fuel consumption, kg/h..... 0.65  
 Composition of mixture oil to fuel ratio, ..... 1:25  
 Temperatures, °C  
     ambient and intake-air..... 40±2  
     crankcase..... 100±2  
     cylinder head..... 220±2  
     cylinder, on Generator Side..... 170±2  
     cylinder, on Starter side..... 195±2  
     exhaust gases..... 350±5  
     Exhaust back pressure, mm of water..... 70±10

The oils were rated on the basis of a number of indicators that characterize their use properties (Table 1).

Table 1. Results of Tests of Additives with AS-9.5 Oil by IM2MD-1ORG Method

Additive (base oil)	Composition, %	Cylinders burnt hours	Cylinders burnt hours	Kerosene consumption, l			F	Kerosene consumption, l			E	Consumption of kerosene burnt in cylinder head
				G	H	I		G	H	I		
AS-9.5 (gasoline)	0	14	5	0.36	0.09	0.16	0.11	0.72				
ACN-300K	15	10.3C	5	0.25	0.13	0.29	0.23	0.98				
CHIN (C <sub>10</sub> -C <sub>12</sub> )	15	8.2C	5	0.38	0.03	0.07	0.05	0.46				
CHIN (C <sub>10</sub> -C <sub>12</sub> )	15	9.9	6	0.3	0.03	0.07	0.05	0.6				
ANR	15	5.7C	5	0.15	0.02	0.03	0.04	0.28				
CHIN (C <sub>10</sub> -C <sub>12</sub> )	15	5.2	1	0.10	0.01	0.12	0.15	0.47				
CHIN (C <sub>10</sub> -C <sub>12</sub> )	15	1	0.2	0.01	0.02	0.07	0.05	0.15				
THIO	15	3.65	1	0.12	0.07	0.15						
HDS-200A	15	0.01	0.3	0.09	0.07	0.12						

Notes: (a) additives; (b) adhesive content, %; (c) deposit index, points; (d) piston-ring mobility; (e) color of deposits on piston, points; (f) amount of deposits on piston, points; (g) on piston heads; (h) on piston heads; (i) on cylinder head; (j) in exhaust port; (k) total; (l) AS-9.5 oil; (m) first ring burned on 360<sup>th</sup> second ring; (n) F-20-0.5 + 3% VNI NP-310 + 1% PMS-Ya + 0.5% EnzhK; (o) all rings free; (p) EnzhK = fatty acids; (q) EnzhK (synthetic fatty acids); (r) same; (s) TPK (triethoxy phosphat); (t) TPK (triethoxy phosphat).

KEY: (a) products; (b) composition of fuel mixture; (c) deposits index; (d) piston-ring mobility; (e) color of deposits on piston, points; (f) amount of deposits on piston heads; (g) on piston heads; (h) on piston heads; (i) on cylinder heads; (j) in exhaust port; (k) total; (l) AS-9.5 oil; (m) first ring burned on 360<sup>th</sup> second ring; (n) F-20-0.5 + 3% VNI NP-310 + 1% PMS-Ya + 0.5% EnzhK; (o) all rings free; (p) EnzhK = fatty acids; (q) EnzhK (synthetic fatty acids); (r) same; (s) TPK (triethoxy phosphat); (t) TPK (triethoxy phosphat).

Table 2. Results of IM2MD-1ORG-Method Tests of Oil Specimens with Domestic and Foreign Additives

Additive	Type	Consumption of kerosene burnt in cylinder head	Consumption of kerosene burnt in cylinder head	Consumption of kerosene burnt in cylinder head			M	N	O	P	Q	R	S	T	U	V	W	X	Y	Z
				E	F	G														
AC-9.5 Gas specimen	1	13.4	13.4	1.23	0.48	0.41	0.36	0.21	0.67											
1. Dioxane	1	10.9	10.9	1.15	0.47	0.40	0.35	0.20	0.52											
2. Dioxane	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
3. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
4. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
5. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
6. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
7. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
8. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
9. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
10. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
11. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
12. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
13. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
14. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
15. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
16. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
17. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
18. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
19. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
20. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
21. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
22. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
23. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
24. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
25. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
26. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
27. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
28. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
29. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
30. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
31. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
32. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
33. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
34. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
35. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
36. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
37. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
38. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
39. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
40. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
41. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
42. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
43. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
44. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
45. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
46. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
47. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
48. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
49. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
50. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
51. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
52. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
53. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
54. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
55. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
56. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
57. Acetone	1	10.7	10.7	1.15	0.47	0.40	0.35	0.20	0.52											
58. Acetone	1	10.7	10.7</																	

[Key to Table 2; except'd.] (p) AS-9.5 + 3.5% VNII NP-360 +  
+ 1% AZNII-TSTATIM; (q) first ring burned on 180°,  
second ring free; (r) AS-9.5 + 3% GFR [East Germany]  
additive for two-stroke gasoline engines; (s) AS-9.5 +  
+ 2% Amoco 661 additive (USA); (t) same; (u) AS-9.5 +  
+ 3% Ecocube 120 additive (USA); (v) AS-9.5 + 1.5% ASK +  
+ 1% SZHK (C<sub>10</sub>-C<sub>18</sub>) + 0.001% PMS-200A; (w) AS-9.5 + 1.5%  
ASK + 1.5% EEZhK + 0.5% FEA (dissubstituted) + 0.001%  
PMS-200A; (x) AS-9.5 + 17.5% ASK + 1% SZHK  
(C<sub>10</sub>-C<sub>18</sub>) + 0.25% VNII NP-354 + 0.001% PMS-200A; (y)  
AS-9.5 + 17.5% T-1 + 1.5% ASK + 1.5% EEZhK + 0.25% VNII  
NP-354 + 0.001% PMS-200A.

Each specimen was tested at least twice; here the disagreement between the results should not exceed +10% for the wear and total-deposits indices, or +0.5 point for the deposit color. The amount of scale is determined to within 0.001 g.

The work of selecting oil additives was begun with study of the functional properties of the individual additives on the bench engine.

It was established that the smallest amount of deposits in the combustion chamber and exhaust ports results from use of the low-ash alkylsulfonate additive ASK, 1.5% by weight of which eliminates piston-ring burning, reduces piston varnishing from 5 to 1 point, and simultaneously increases the amount of deposits on the piston head, in the cylinder head, and in the exhaust port (see Table 1).

Fatty acid ethyl ester (EEZhK) and synthetic fatty acids (SZHK) reduce the amount of deposits substantially but do not eliminate burning-on of the piston rings and do not reduce varnishing of the piston. More than 1% of SZHK in the oil causes deterioration of all motor-property indicators. Combined with ASK and SZHK, PMS-200A additive improves all indicators.

The studies established that each of the additives tested does not, taken alone, improve the properties of the base oil in all of the rating indicators. Subsequent development work on oils for two-stroke gasoline engines was directed toward adjustment of the additive combination.

Table 2 presents test results for mixes developed on the basis of ASK additive in combination with EEZhK, SZHK, and PMS-200A, for comparison with AS-9.5 base oil and with foreign additives. Mixes based on ASK are more effective. Additive mixes containing synthetic fatty acids (C<sub>10</sub>-C<sub>18</sub>) fraction, which were used for 650-1000 hours in Druzhba-4 gasoline-powered saws, produced somewhat better results than mixes of the same additive containing EEZhK. When AS-9.5 oil was used with an additive mix consisting of 1.5% ASK, 1% SZHK, and 0.001% PMS-200A, the service life of the Druzhba-4 engine was increased by a factor of 2. This oil is recommended by the Interdepartmental commission for experimental use. The good use properties of AS-10 oil with

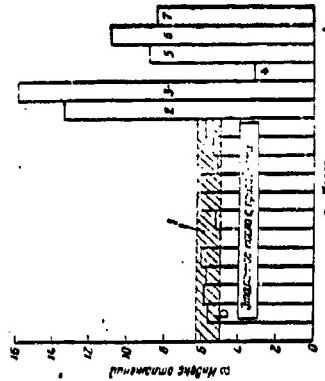


FIG. 1. Reproducibility of test results and comparison of various oils on the basis of the "deposits index" indicator. 1) scattering region of control-oil test results; 2) AS-9.5 (without additive); 3) AK-15 (without additive); 4) VNII MP mix; 5) AS-9.5 + 3% Ecocube 120; 6) AS-9.5 + 3% GDR additive; 7) AS-9.5 + 2.5% Amoco 661.  
KEY: (a) deposits index; (b) control oil with additives; (c) oil.

17.5% of T-1 fuel, 1.5% ASK, 1% SZHK, 0.25% VNII MP-154, and 0.001% PMS-200A and a similar mix with 1.5% EEZhK were confirmed by the results of use tests extending over the entire lifetime of SD-60B engines. In tests of engine performance on AS-9.5 oil without additives (mixture composition 1:25), it was necessary to replace the piston pin and the connecting-rod top-end insert after 660 hours because of wear. Engines ran for more than 700 hours on AS-9.5 oil with the additive corporation developed (mixture composition 1:33) without replacement of parts or removal of deposits.

The method developed makes it possible to obtain a reproducibility of the results that is adequate for motor tests, along with sharp enough differentiation between different oil qualities (Fig. 1).

The installation developed performs reliably over a broad cylinder-temperature range. OILS can be rated in research studies at cylinder temperatures from 170 to 230°C. The maximum cylinder temperature for tests of various oils and additives (Fig. 1).

### CONCLUSIONS

1. A method was developed for motor rating of the use properties of oils made for two-stroke gasoline engines.
2. The method can be used to rate oils with and without additives, to determine the functional properties of specific additives, and to adjust optimum additive mixes.
3. The method yields satisfactory reproducibility in the test installation. The divergence of two parallel tests is  $\pm 10\%$  for the varnishing points index and the total amount of deposits.
4. Five tenth installation and the method elaborated for it have enabled the VNIIT MP to develop special two-stroke gasoline engines whose operational characteristics are considerably better than those of the commercial oils currently being produced.
5. The motor-rating method can be recommended for State Standard and Departmental Technical Specifications, at petroleum refineries, by scientific-research organizations, at petroleum refineries, and by consumers to check oil quality.

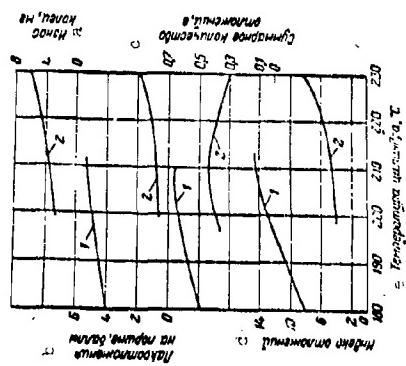


Fig. 2. Influence of thermal load of IM-210FG bench engine on test results for various oils. 1) PS-11 oil (without additive); 2) Fr-II oil with 1.5% ASK, 1% SK, and 0.001% PMS-200A.  
KEY: (a) ring wear, mm; (b) piston varnishing points; (c) total amount of deposits, g; (d) deposits index; (e) cylinder temperature, °C.

ring wear additives, it is usually 180–200°C. In ten-hour sequential tests of the oil, its anti-wear properties can be rated quite reliably within 40–50 hours. When oil was used without additive, in 40 hours of operation at cylinder temperatures between 160 and 210°C, the total piston环 wear was 0.027 mm but only 0.017 mm with 1.5% ASK, 1% SK, and 0.001% PMS-200A over the same running time at cylinder temperatures of 200–250°C.

A full-scale bench installation has now been built around a 1.0-litre engine and can be used to test oils for two-stroke gasoline engines.

Footnote

77 <sup>1</sup>A stationary single-cylinder two-stroke engine with two-channel return-counterflow scavenging.

IN-STORAGE PROPERTY CHANGES OF OILS OF VARIOUS CLASSES AND DETERMINATION OF THEIR PURITY

CHANGE IN THERMAL-OXIDATION STABILITY AND DETERGENT POTENTIAL OF ADDITIVE OILS DURING STORAGE

K.K. Papok and B.S. Zuseva

With the object of determining how the thermal-oxidation stabilities and detergent potentials of additive oils change during storage, oils of various classes (from Premium to Series 3) were prepared with DS-11 oil as a base and with various combinations of foreign and domestic additives and placed in long-term storage.

A consignment of oils with foreign additives was prepared under laboratory conditions in August 1961. Each specimen had a volume of 0.5 liter. The oil samples were stored in glass bottles in diffuse light in a closed unheated warehouse. Control analyses were made in February 1964 and April 1965.

The analyses (Table 1) showed that after 2 years and 7 months of storage, all oils with mixes of foreign barium and calcium additives retained thermal-oxidation stability (except for the calcium-additive Series 3 oil), while after 3 years and 9 months of storage, thermal-oxidation stability had declined by 10-25 minutes in 50% of the specimens, namely, in 115 of the Heavy Duty, Series 1 and Series 2 classes with tarlike additives and in Premium, Series 1 and Series 3 oils with the calcium additives.

Table 1. Change in Properties of Various Classes of Oils During Storage in Glass Bottles

a. Номер	b. Содержание стабилизатора в масле ДС-11	Термостабильность, часы с способностью к окислению						c. Массовая концентрация, %
		d. Оригинальная	e. Стабилизированная	f. Стабилизированная с добавкой антиоксиданта	g. Стабилизированная с добавкой бария	h. Стабилизированная с добавкой бария + 0.7% антиоксиданта	i. Стабилизированная с добавкой бария + 2.6% кальция	
1. Примул	0.7% антиоксидантный	52	49	55	10	5	10	
2. Хей-Дьюти	0.7% баритовый	60	56	50	20	20	60	
3. Серия 1	0.25% баритовый + 4% антиоксидантный	60	53	34	50	50	-	
4. Серия 2	0.7% баритовый + 4% антиоксидантный	95	84	-	75	85	-	
5. Серия 3	1.5% баритовый + 4% антиоксидантный	76	73	-	73	85	-	
6. Примул	0.6% антиоксидантный	45	45	35	20	5	5	
7. Хей-Дьюти	0.6% антиоксидантный + 2.6% кальциевый	45	40	45	60	30	35	
8. Серия 1	0.5% антиоксидантный + 4.3% кальциевый	45	40	45	35	65	-	
9. Серия 2	1.0% кальциевый	47	46	45	-	50	-	
10. Серия 3	1.5% кальциевый	68	52	55	-	50	-	
11. Для судов морской плаваги	25% кальциевый	110	92	85	-	90	-	

KEY: (a) oil; (b) content of additives in DS-11 oil; (c) thermal-oxidation stability, minutes; (d) original specimen; (e) after 2 years; (f) after 5 years; (g) after 1 year; (h) detergent potential; (i) antioxidant potential; (j) Premium; (k) Heavy Duty; (l) C-4 antifouling + 0.7% barium; (m) C-4 antifouling + 0.7% barium + 2.6% calcium; (n) ...% calcium; (p) for marine engines.

Detergent potential remained unchanged in all oils after three years and nine months of storage, except for the Premium and Heavy Duty oils with calcium additives, which showed a sharp drop in detergent potential even after 2 years and 7 months.

In August 1962, a consignment of oil with various additives, prepared under semiindustrial conditions and packaged in steel drums, was obtained from the NII RP. 0.5-liter samples of the oils were drawn from these drums into glass bottles and placed in a closed unheated warehouse for storage. The drums were then stored on an outdoor platform.

Control analyses, the results of which appear in Table 2, were made in February 1964 and April 1965.

As we see, even after 1.5 years of storage, whether in the glass bottles or in the steel drums, the thermal-oxidation stabilities of all the oils except the Series 1 and Series 2 oils with calcium additives had dropped by 12-30 min, and after 2

Table 2. Change in Properties of Oils of Various Classes During Storage\*

a. Номер	b. Стабилизатор	Стабилизация масла при хранении						c. Массовая концентрация, %
		d. Оригинальная	e. Стабилизированная	f. Стабилизированная с добавкой антиоксиданта	g. Стабилизированная с добавкой бария	h. Стабилизированная с добавкой бария + 0.7% антиоксиданта	i. Стабилизированная с добавкой бария + 2.6% кальция	
1. Примул	Баритовый	55	42	36	28	20	10	
2. Хей-Дьюти	Баритовый	62	46	30	29	20	20	
3. Серия 1	Баритовый	62	47	39	30	20	20	
4. Серия 2	Баритовый	75	50	40	35	25	20	
5. Серия 3	Баритовый	70	41	35	35	25	20	
6. С-4 антифулинг	Баритовый	48	32	28	20	20	20	
7. Хей-Дьюти	Баритовый	45	32	28	20	20	20	
8. Серия 1	Баритовый	35	24	20	20	20	20	
9. Серия 2	Баритовый	40	32	39	30	20	20	
10. Серия 3	Баритовый	75	64	55	55	25	20	

\*The additive contents of the various oils are the same as in Table 1.

KEY: (a) oil; (b) storage conditions; (c) original specimen; (d) thermal-oxidation stability, minutes; (e) after 1.5 years of storage; (f) after 2 years; (g) after 5 years; (h) detergent potential; (i) Premium; (j) Heavy Duty; (k) with barium; (l) with calcium; (m) Series...; (n) with calcium additives.

years and 6 months of storage, a further decrease in thermal-oxidation stability was noted in the oil stored in glass bottles, while further observations were made on the oil in the drums after 1.5 years.

Except for Premium oil with barium additives, the detergent potential of all oils had undergone no change after 1.5 years of storage, while after 2 years and 8 months, all oils with calcium additives showed a 20-60-unit loss of detergent potential.

It is interesting to note that the oils prepared in the laboratory (August 1961) were more stable during storage. In glass bottles prepared under semiindustrial conditions (see Tables 1 and 2). This provides a basis for the assumption that the conditions of mixing of the oils with the additives influence storage stability.

The results of analyses of the oils with Soviet additives appear in Table 3.

Table 3. In-Storage Property Changes of Oils of Various Classes With Soviet Additives

a Series	b Catalytic stabilizer in Series RC-11	c Purified specimens	d Original specimens	Type, liters		e f g h i j k l m n o p q r s t u v w x y z	Liquor number	Weight mg		
				original specimen	specimen after 1 year storage					
1 Series 1	J 3% BHIIH HN-370+1% PMS-Ya + 0.5% JZ-23k + 0.005% HNC-200A	K Catalyst tape 1B Gorok Gorok	K Catalyst tape 1B Gorok Gorok	65	62	45	30	30	40	-
m Series 1	n 5% BHIIH HN-370+2% PMS-Ya + 0.5% JZ-23k + 0.005% HNC-200A	K Catalyst tape 1B Gorok Gorok	K Catalyst tape 1B Gorok Gorok	65	65	-	30	30	40	-
m Series 2	O 4% BHIIH HN-370+4% PMS-Ya + 0.5% JZ-23k + 0.005% HNC-200A	K Catalyst tape 1B Gorok Gorok	K Catalyst tape 1B Gorok Gorok	77	62	62	30*	30*	0	-
m Series 3	P 15% BHIIH HN-370+6% PMS-Ya + 0.5% JZ-23k + 0.005% HNC-200A	K Catalyst tape 1B Gorok Gorok	K Catalyst tape 1B Gorok Gorok	75	75	-	30*	30*	0	-
				69	62	30	30	30	0	-
				69	62	-	30*	30*	0	-

\*Filtered specimens.

KEY: (a) oil; (b) additive content in DS-11 oil; (c) storage conditions; (d) thermal-oxidation stability, minutes; (e) original specimen; (f) after 1.5 years of storage; (g) after 2 years; (h) months of storage; (i) detergent potential, %; (j) Heavy Duty; (k) 3% VNII NP-370 + 1% PMS-Ya + 0.5% LZ-23k + 0.005% PMS-200A; (l) in steel drums; (m) Series 1; (n) 5% VNII NP-370 + 2% PMS-Ya + 0.5% LZ-23k + 0.005% PMS-200A; (o) 11% VNII NP-370 + 4% PMS-Ya + 0.5% LZ-23k + 0.005% PMS-200A; (p) 15% VNII NP-370 + 6% PMS-Ya + 0.5% LZ-23k + 0.005% PMS-200A.

After 1.5 years of storage, the thermal-oxidation stability of all oils (except for two specimens) remained practically unchanged, but after 2 years and 8 months of storage in glass, they fell by 8-20 min. The detergent potentials of the Heavy Duty and Series 1 oils had not changed after 2 years and 8 months of storage; for the Series 2 and Series 3 oils, they had fallen by 20-55 units after 1.5 years of storage, and to zero after 2 years and 8 months.

#### INVESTIGATION OF PURITY OF MOTOR OILS AND ADDITIVES

The following method was worked out for determination of the purity of the oils and additives. The oil or additive specimen to be studied was dissolved in Galosha gasoline at room temperature and passed under vacuum (residual pressure 20-30 mm Hg) through a No. 4 biological filter 27 mm in diameter, which was placed in a two-part metal funnel (GOST [All-Union State Standard] 10734-64). To determine the purity of the oils, 5-gram weighed specimens were taken and dissolved in 45 ml of Galosha gasoline; to determine additive purity, 1-gram weighed samples were dissolved in 49 ml of Galosha gasoline.

The purity of an oil or additive is determined from the filtration number and the amount of dirt in the oil. The filtration number is determined from the number of filters needed for complete filtering of the entire oil or additive solution with filter change mandatory at 5-minute intervals.

The amount of dirt is determined from the difference in the weights of the biological filter before and after filtration of the oil solution (bringing it to constant weight at 100°C). When it is necessary to change filters several times, as when the products being tested contain large amounts of contamination, the total amount of deposits on all of the filters is taken as the fouling index of the oil. The amount of dirt is expressed in mg/100 g of the oil or additive being tested.

The disagreement between parallel experiments may not exceed 200 mg/100 g for dirt contents up to 200 mg/100 g, or 10% of the smallest result for dirt contents above 200 mg/100 g. The fouling of the oil can also be rated tentatively by visual inspection on the basis of filter color.

It was found on checking the purity of the motor oils without additives that in general, all of the commercial oils show high purity: the number of filtrations was 1, i.e., they were thoroughly filtered by a single filter, and the amount of dirt did not exceed 30 mg/100 g.

The purity of motor oils containing various additives fluctuates in a very broad range: as many as 6 filters were required to clear some specimens, and the amount of dirt ranged up to 300 mg/100 g (Table 4).

Oils with the Monto, Santolube, and Orobit additives showed high purity: the filtration number was 1, and the amount of dirt ranged from 26 to 40 mg/100 g. On the other hand, oils with the VNII NP-370, BFK-1, SB-3, and other additives were very dirty: they required up to 6 filtrations and contained up to 308 mg/100 g of dirt.

The purity change of mineral oils under the influence of additives is accounted for by inadequate purity of the additives themselves (Table 5). We see from the table that the filtration

Table 4. Furtiles of Various Classes of Oil

a	b	c	Состав участка		
			d	e	f
<i>Соединение промежуточное в зоне ЛС-44</i>					
Б	Б	Б			
Без	Саранов-633 + 0,7% Марко-613	-			
Без	Орлов-287	-			
Без	СН-11 + 4,2% БФ-11	-			
Без	СН-31 + 3% БФ-11	-			
Без	Малыш-31 + 0,7% БФ-00К	-			
Без	Самаров-633 + 1,5% Марко-613 Б	-			
Без	Орлов-287 + 0,7% Марко-285	-			
Без	ВИИ-370 + 1% Марко-613 + 0,5% ЛС-23 + 0,05%	ПЧС-20ДА	2		
Без	БФ-1 + 5% СН-3 + 0,05%	ПЧС-20ДА	2		
Без	ЛАНН-31 + 1,5% БФ-1	-			
Без	Саранов-633 + 4% Марко-613	-			
Без	Орлов-287 + 3,5% Марко-613	ПЧС-20ДА	2		
Без	ВИИ-370 + 1% Марко-613 + 0,5% ЛС-23 + 0,05%	ПЧС-20ДА	2		
Без	БФ-1 + 3% СН-3 + 0,05%	ЛС-23 + 0,05%	2		
Без	ПЧС-20ДА	-	2		
У	Без	БФ-1			
У	Без	ЛАНН-317 + 4% БФ-1			
Х	Без	Саранов-633 + 9% Марко-613			
Х	Без	Орлов-287			
У	Без	ВИИ-370 + 0,5% Марко-613 + 0,5% ЛС-23 + 0,05%	ПЧС-20ДА	6	
У	Без	БФ-1 + 4% СН-3 + 0,5% ЛС-23 + 0,05%	ПЧС-20ДА	2	
аа	Без	БФ-1 + 4% СН-3 + 0,05%	ПЧС-20ДА	2	
аа	Без	БФ-1 + 4% СН-3 + 0,05%	ПЧС-20ДА	2	
3	Без	Марко-20ДА			
3	Без	Орлов-287			
cc	Без	БФ-212А			
dd	Без	ПЧС-20ДА			
ee	Без	ПЧС-20ДА			
ff	Без	ПЧС-20ДА			

number was 1 for the LAN-217, Santolube-493, Monto-613 and 702 and Orchis-267, 205 $\mu$ , and 212A additives, while they contained no more than 80 mg./1.00 of dry.

Table 5. Purities of Various Additives

Группа	Номер	Состав		Код	Причина	Место	Время
		С	Г				
Санитар-6-33	Чонго-613						
	Очог-154						
	Ланко-702						
	Ланко-317						
	Ланко-335						
	Бон-4						
	Бон-22						
	ВИИИ НИ-371						
	ВИИИ НИ-370						
	АНИИ-ДИНАМ-1						
	АНИИ-7						
	ВИИИ НИ-22						
	ВИИИ НИ-370						
	ВИИИ НИ-457						
	ГБ-3						
	ПМС-2						

G	Sorta
3	100% Могилев-202 бб
4	100% Орловец-212а
5	100% БИШ-1Н-370+5% ПМС-ЧИ+0,5% НС-23к+
6	100% БИШ-1Н-370+5% ПМС-ХВА
7	100% БЕ-1+6% СЕ-3+0,5% НС-23к+0,065%
8	100% ПМС-ЗАМДА
9	95% ДАННИ-317+10% БФК-1+0,005% НС-200А
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However, most of the additives have indicators inferior to those noted above, and additives such as PMS<sup>1</sup>, Ya, SB-3, and VNJI SP-370 were very dirty (from 850 to 2070 mg/100g) and had filtration numbers from 4 to 9. However, any of the additives may be prepared in different purities.

The purities of laboratory-mixed (August 1961) and semi-industrial (April 1964) oils with foreign additives and domestic semindustrially-mixed oils were checked. As we see from Table 6, the semindustrially mixed oils are considerably dirtier than the lab-mixed oils, and this is apparently the explanation for the stability differences during storage.

It is recommended that the purity-rating method be used to monitor the purity of oils and additives, using the norms given in Table 7 as a guide.

Table 6. Purities of Oils of Various Classes Mixed in the Laboratory and Under Semilandustrial Conditions

**KEY:** (a) oil; (b) purity; (c) number of filtrations; (d) laboratory mixing; (e) semiindustrial mixing; (f) dirt, mg/100G; (g) with imported barium additives; (h) Premium; (i) Heavy Duty; (j) Series 33; (l) imported calcium additives; (m) with domestic YNT NP additives.

Table 7. Acceptable purity standards

	D	E	F	G	H	I	J
A	Примечание	С. оценка затраты	Д. оценка затраты	И. оценка затраты	К. оценка затраты	Л. оценка затраты	М. оценка затраты

**KEY:** (a) Product; (b) Purity; (c) number of filtrations; (d) dirt, mg/100S (not above); (e) oil without additives; (f) oil with additives; (g) additives.

## CONCLUSIONS

1. During prolonged storage, thermal-oxidation stability and detergent potential decline in oils of various classes (from Premium to Series 3, inclusive) with both foreign and domestic additives.

Inhibitors that terminate chains without consumption (in the first result) or the NH<sub>2</sub> group, but with participation of the phenol OH group [2].

According to the second view of the synergistic effect, inhibitors are classified into three kinetic groups that differ in chemical structure and are capable of deactivating various intermediate products on which initiation and development of the chain oxidation reaction depend [3-5]. It has been established by research at the AllUnion Institute of Heat Engineering (BTU) that Group I retarders interact only with free hydrocarbon radicals R·, and not with hydroperoxides ROOH or free peroxide radicals RO<sub>2</sub>·; Group II retarders react vigorously with hydroperoxides and do not react with RO<sub>2</sub>·, and do not react with hydrocarbon radicals R·; Group III retarders react with R· and RO<sub>2</sub>· and interact either sluggishly or not at all with hydroperoxides [5].

In this context, we may assume that in the simultaneous presence of antioxidants of various groups (I and II, II and III, I and III), their effects formed during the initiation phase, while those that still exist and have time to convert to RO<sub>2</sub>·, are handled by the group II antioxidant.

For example, we might assume that in the combined presence of group I and II retarders, the group II antioxidant deactivate most of the R· radicals formed during the initiation phase, while those that still exist and have time to convert to RO<sub>2</sub>·, are handled by the group II antioxidant.

Inhibitors of Groups II and III (to a certain degree III more, however, produce a synergistic effect by interacting with some unreacted hydroperoxides, thus preventing branching of the reaction chain.

The first hypothesis in conjunction with the fact that inhibiting agents are capable in many cases of forming complexes with hydroperoxides containing conjugated bonds, e.g., in quinones, etc., has been substantiated on the example of primary aromatic amines and some substituted phenols. These complexed compounds exhibit properties, however, that theory does not always predict. In addition, the action of aromatic inhibitors (e.g., 4-aminophenol, 4-aminobiphenol, 4-aminonaphthalene, etc.) is not due to the participation of hydroperoxides RO<sub>2</sub>·, but rather to the participation of hydrocarbon radicals R·, alkoy radicals RO·, etc. We believe this theory correctly characterizes the experimentally established differences between the abilities of different inhibitors to retard the oxidation process, i.e., to react with different intermediate products of the oxidation process.

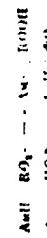
The second theory takes these factors into account. In particular, the selection of a broad range of inhibitor mixtures can reinforce the synergistic effect. The possibility of such a selection has been confirmed experimentally under laboratory and natural conditions.

## USE OF PAIRED ANTIOXIDANTS FOR TURBINE OILS

K.I. Ivanchy, Ye.F. Vilyamzaya, A.A. Buzzetskly  
and A.N. Al'ebendrov

There is as yet no generally accepted theory that explains the phenomenon observed when mixtures of hydrocarbon-extraction inhibitors (antioxidants, retarders) are used, i.e., the mutual enhancement of their action that is observed in many cases.

In this present article, we shall try to explain this phenomenon. As a rule, one of them (in the case of aromatic amines, for example) acts as a retarder, while the other (e.g., a phenol, etc.) acts as an antioxidant. In the first case, the formation of hydroperoxides is inhibited, and in the second, the propagation of the reaction is retarded. In this manner, the two types of retarders act in concert, and the overall inhibition is increased. This is the case, for example, in the interaction of 4-aminonaphthalene with 4-aminophenol, 4-aminobiphenol, 4-aminonaphthalene, etc.



After the orientation to para-aminodiphenyl [1], or as a result of formation of condensate of 4-aminobiphenol with the amine and the generation of hydroxy-constituting products produced with a hydrocarbon at the nitrogen:  $\text{H}-\text{N}-\text{H}\cdots\text{HO}-$ , which are active

Laboratory and use tests have shown that given proper selection, the stabilizing action of two additives on turbine oils exceeds that of either of them taken alone at twice the concentration. This proves that we are dealing not with simple addition of the effects of two antioxidants, but with mutual enhancement of their activities. It has also been proven that a mixture of antioxidants of the same kinetic group does not, as a rule, produce the synergistic effect [6].

#### Effectiveness of Paired Antioxidants Under Laboratory Conditions

a. Reagent	Oils after 100 hours at 150°C			
	b C <sub>18</sub> H <sub>34</sub>	c C <sub>18</sub> H <sub>34</sub> + KOH/g	d C <sub>18</sub> H <sub>34</sub> + KOH/g + C <sub>18</sub> H <sub>34</sub>	e C <sub>18</sub> H <sub>34</sub> + KOH/g + C <sub>18</sub> H <sub>34</sub> + traces
Machine oil I	0.61	0.48	0.48	0.48
To ace + 0.02% component A	0.78	0.66	0.66	0.66
o + 0.1% component B	0.17	0.23	0.23	0.23
o + 0.01% component A	0.04	0.04	0.04	0.04
o + 0.03% component B	0.05	0.05	0.05	0.05
Machine oil II	0.48	0.38	0.38	0.38
To ace + 0.01% component A	0.14	0	0	0
o + 0.05% component B	0.25	0.04	0.04	0.04
o + 0.01% component A + 0.03% component B	0.07	0.07	0.07	0.07
o + 0.03% component B + 0.01% component A	0	0	0	0

KEY: (a) product; (b) over-all stability according to GOST 981-55; (c) acid number, mg of KOH/g; (d) sediment content, %; (e) turbine oil I; (f) same + ...% of component A; (g) same + ...% of component B; (h) traces; (1) turbine oil II.

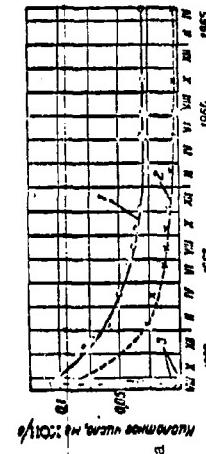


Fig. 1. Change in acid number of oil after introduction of paired antioxidant. 1) total acid number; 2) acid number due to water-soluble acids; 3) introduction of paired antioxidant.  
KEY: (a) acid number, mg of KOH/g; (b) test time.

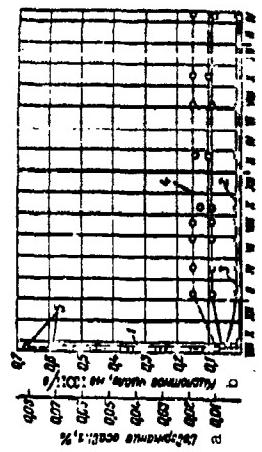


Fig. 2. Change in stability of oil after introduction of paired antioxidant. 1) acid number; 2) content of nonvolatile acids; 3) content of volatile acids; 4) sediment content; 5) introduction of paired antioxidant.  
KEY: (a) sediment content, %; (b) acid number, mg of KOH/g; (c) test time.

Of special practical interest for inhibiting oxidative aging of lubricating and insulating oils as a result of reaction of antioxidants and synergistic effects is the use of mixtures of antioxidants (i.e., substances in which, in addition to the true inhibitors, there are re-arrangers, substances that terminate reaction chains), there are retarders, with other mechanisms of action - primarily deactivators and passivators for metals (chiefly copper and iron) that come into contact with turbine and transformer oils under use conditions, and catalytically accelerate their aging.

It must be remembered in testing various antioxidant pairs that they must dissolve well in oils and ensure stability without detriment to other use properties.

A paired antioxidant consisting of 0.1% 8-hydroxy-β-naphthylamine and 0.03% phenyl-β-naphthylamine, a Group I antioxidant, was tested after laboratory study in the oil system of a hydrocracker unit generator. The oil with this antioxidant pair was used for 1.5 years without the use of adsorbents. After 1.5 years, the acid number of the stabilized oil was considerably lower than those of unstabilized oils previously used in this unit. The adsorbents for the same period of time.

Tests were run on one more mixture, which consists of 0.1% Group III antioxidant (component A) that has similar physical properties of a metal deactivator and a Group I antioxidant (component B). The effectiveness of this mixture as

under laboratory conditions according to GOST 981-55. The test results are given in the table.

The table shows that the proposed antioxidant pair has a very sharp inhibiting effect on the aging of turbine oil. The acid number after aging was smaller by a factor of 12 than that of the pure oil, and smaller by a factor of 3 than the acid number of the same oil with twice the concentration of one of the components. The amount of sediment is also smaller, in confirmation of the effectiveness of this paired antioxidant.

Use tests were then run with this antioxidant pair in a 24,000-kW turbogenerator.

Figure 1 shows the change in the acid number of the stabilized oil during operation of the turbogenerator. Figure 2 shows the change in its stability (after GOST 981-55) after introduction of the paired antioxidant.

It was established by the tests that after 3.5 years of use, the oil had an acid number of 0.03 mg of KOH/g, i.e., it was practically the same as the norm for fresh oil (0.02 mg of KOH/g). It should be noted that no additive was introduced into the fresh oil over the 3.5 years. It was established on inspection of the turbine (after 2 years of operation), that its oil system did not require cleaning. Operating experience from previous years shows that without the paired antioxidant, oil lasted 2 years in this turbine only with continuous adsorbent regeneration.

The adsorbent had to be changed repeatedly, with the resulting higher labor cost and a considerable increase in use of the expensive adsorbent.

The same paired antioxidant was tested in another turbine. Figure 3 shows the changes of the oil's indicators in this turbine. The tests are being continued successfully. The use tests show that the proposed paired antioxidant inhibits the turbine-oil aging process more actively than other antioxidant additives and does not require special checking by servicing personnel. At this writing, use tests are being run with the paired antioxidant in four turbogenerators.

#### CONCLUSIONS

1. Mixtures of antioxidants for turbine oils, adjusted on the basis of theoretical notions developed by the authors, give a pronounced synergistic effect when tested under laboratory conditions.

2. The results of the laboratory tests are confirmed by data from use testing of the paired antioxidant, in the absence of which the original properties of the oil are retained fully over a long operating interval.

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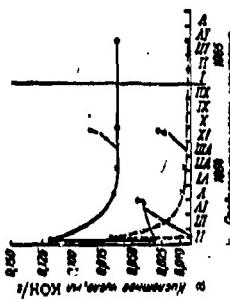


FIG. 3. Change in acid number of oil after introduction of paired antioxidant. 1) total acid number; 2) acid number due to water-soluble acids; 3) introduction of paired antioxidant. KEY: (a) acid number, mg of KOH/g; (b) test time.

Basic Physicochemical Indicators of Transformer Oils

a name	b Oxidants	Copper Oxidized and 2,6-d <sup>-tert</sup> -butyl-4-phenyl-						Copper Oxidized and 2,6-d <sup>-tert</sup> -butyl-4-phenyl-		
		c 100°C	d 150°C	e 180°C	f 200°C	g 220°C	h 240°C	i 260°C	j 280°C	k 300°C
b. <i>nickel</i> <i>nitrate</i>	tinocene- magnesium oxide	0.0872	1.0752	27.2	1.1	11.3	42.7	6.12	1.72	
c. <i>nickel</i> <i>nitrate</i>	Aluminum oxide	0.0839	1.0532	25.8	0.4	14.2	38.3	6.14	1.53	
d. <i>nickel</i> <i>nitrate</i>	Kuchino- magnesium oxide	0.0857	1.0517	22.1	2.5	13.3	45.1	6.18	1.70	
e. <i>nickel</i> <i>nitrate</i>	Tin oxide	0.0867	1.0572	19.6	2.5	22.2	38.0	6.56	1.48	
f. <i>nickel</i> <i>nitrate</i>	Quaternary amine	0.0610	0.4720	26.7	0.3	5.1	31.6	0.14	1.65	
g. <i>nickel</i> <i>nitrate</i>	Terpineol	0.0870	1.0564	25.3	0.2	30.6	29.5	0.72	1.13	
h. <i>nickel</i> <i>nitrate</i>	Tartaric anhydride	0.0914	1.0526	25.9	0.7	15.3	42.8	0.12	1.61	
i. <i>nickel</i> <i>nitrate</i>	Styrene	0.0914	1.0526	25.9	0.7	15.3	42.8	0.12	1.61	

KEY: (a) salt; (b) organic; (c) viscosity; (d) reactivity; (e) reactivity at 100°C; (f) tin; (g) tin; (h) from mixed BAKI; (i) structural group analysis; (j) from mixed BAKI; (k) adsorption; (l) from mixed BAKI; (m) a zinc base; (n) from fraction V; (o) salt; (p) tin; (q) from Tugayev crude; (r) phenol; (s) from aromatic hydrocarbons; (t) impurities; (u) contact refining; (v) impurities.

Passivators act by a somewhat different mechanism than the additives. They form a catalytically inactive film on the surface. It is assumed that the passivator is a tenth of a micrometer thick.

The passivating additives include certain amines [2,4], quite often, the same compound acts as both a deactivator and a passivator [1].

Reference [3] describes experiments in which additives of this type were used separately and in combinations, but it would be interesting to study the manner in which these additives increase the stability of domestic commercial transformer oils of various origins (Table).

The additives disalicylideneethylenediamine (a deactivator) and anthranilic acid (a passivator) were studied, along with combinations of these additives with one another and with anti-oxidants: phenyl-B-naphthylamine and 2,6-d<sup>-tert</sup>-butyl-4-phenyl-phenol (lonol).

The effectiveness of these additives was rated by laboratory methods, which simulate the most important conditions to which the oil is subjected in transformers. One of the methods consists in oxidizing the oil for 44 hours with oxygen under static

#### ADITIVE MIXES FOR STABILIZATION OF TRANSFORMER OILS

M. S. Shakhovitch

Use of oil-stabilizing additives prolongs the useful lives of transformers, improves their reliability, and permits improvements to their design. Among the insulating and structural materials used in the transformer, copper is the most active participant of the oxidation of materials incalculating oil.

Under use conditions, copper structures come into contact with the oil, and dissolved copper is found in the oil itself, in the form of salts of organic acids.

Used transformer oil contains up to 0.0001% of copper [1]. It has been shown [2] that the amounts of antioxidant additives consumed in inhibiting hydrocarbon oxidation in the presence of copper are considerably larger than under the same conditions without the catalyst. With the large amount of copper present when oil is oxidized in transformers, it is necessary to use very large amounts of antioxidants (0.3-0.5% by weight).

The stability of transformer oils can be increased much more effectively by using deactivating and passivating additives. Deactivators produce their effect by interacting chemically with salts of metals dissolved in the oil. This leads to formation of chelate complexes in which the metal is screened and cannot act as a catalyst. The complexes formed may settle out or remain dissolved in the oil.

Salicyldenes - Schiff bases obtained by condensing amines with aromatic aldehydes - are used as deactivators.

[Key to FIG. 1, cont'd.] without additive;  
 14) same oil with 0.05% anthranilic acid.  
 The tan  $\delta$  values of the oils (at 70°C. in  
 2) after oxidation are given in parentheses.  
 KEY: (a) amount of oxygen absorbed, ml/100 ml  
 of oil; (b) test time, hours.

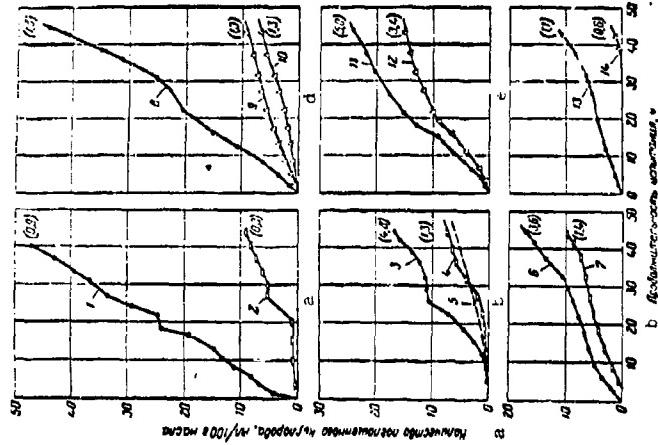


Fig. 1. Kinetic oxygen absorption curves of transformer oils with additives during oxidation in an electric field (a-f): 1) imported oil (GDR) without additive; 2) same oil with 0.05% anthranilic acid; 3) oil from Baku crudes without additive; 4) same oil with 0.05% anthranilic acid; 5) oxidation without copper; 6) oil from Tuyumay crude, hydrorefined and contact-refined without additive; 7) same oil with 0.05% anthranilic acid; 8) oil from Tuyumay crude, phenol refining, without additive; 9) same oil with 0.05% anthranilic acid and 0.3% Ionol; 10) same oil with 0.05% 4-isopropylideneethylenediamine and 0.3% Ionol; 11) oil from Anasias'ev crude without additive; 12) same oil with 0.5% anthranilic acid; 13) oil from Buzovinsk crude, adsorption refining. [Key cont'd. on page 104]

conditions in the presence of catalysts (copper and iron), in an electric field of 49 kV/cm at 100°C. The oxidizability of the oil is evaluated from the rate of oxygen absorption and from the change in the properties of the oil. In another method, the oil is oxidized without an electric field for 1000 hours at 70°C. in the presence of a copper catalyst with air. Oxidation rate for 1000 hours is determined by the change in viscosity. Indicators of the oil after 240, 480 and 720 hours. Together, these laboratory methods yield a preliminary rating of the performance properties of transformer oils with and without additives.

As is now customary in evaluating the quality of transformer oils, the data from the laboratory experiments were compared with results from tests of oil specimens with additive mixes in small bench transformers [5].

It was established by the preliminary experiments that 0.2% by weight of the antioxidant additive Ionol is introduced into transformer oils (see Table 1), stability is not always improved, especially in the case of the [sic] elektropolymer indicators (the tangent of indicator loss angle  $\tan \delta$ ).

The results obtained when deactivating and passivating additives are used are different. Tests in an electric field indicate (Fig. 1) that on addition of 0.05% of anthranilic acid to the various oils, the oxidation process is retarded appreciably. The kinetic curves of oxygen absorption by the oil with the anthranilic acid additive are smooth and coincide almost perfectly with the oxidation curves of the corresponding oil without the additive or catalyst. The oxidation curves of oils with anthranilic acid characterize the slow absorption of oxygen at the beginning of oxidation. Without the extended induction period that is usually observed in oxidation of oils with inhibition additives.

Stability of the indicator tan  $\delta$ , which characterizes the insulating properties of the oils, is highly important for the performance of transformer oils. During oxidation, the tan  $\delta$  of oils with anthranilic acid increases much more slowly than the oils without this additive.

In analyzing the results of 1000-hour oil-oxidation tests (Fig. 2), we note that the stabilizing action of anthranilic acid manifests itself in the oxidation of oil from various crudes containing various amounts of carbon in aromatic rings (2).

the oil together with the passivator. Then, to the extent that copper does not manifest its catalytic activity, inhibitor consumption decreases and the oil becomes more stable.

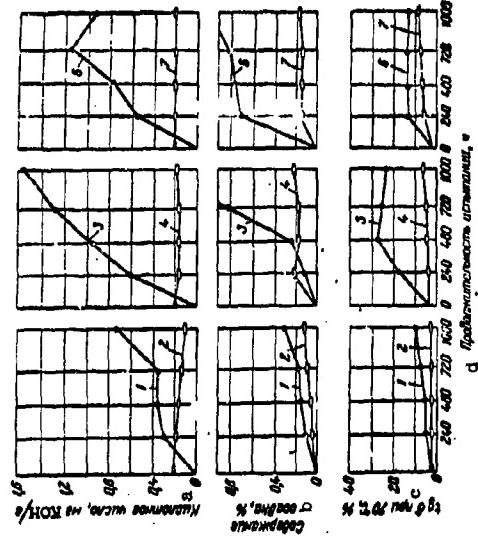


Fig. 2. Kinetic curves of indicator changes during oxidation (1000 h) of transformer oils with 0.05% anthranilic acid (for legend see FIG. 1).  
KEY: (a) acid number, mg of KOH/g; (b) sediment content, %; (c) tan  $\delta$  at 70°C, %; (d) test time, hours.

Oils with anthranilic acid show a typical slow increase in tan  $\delta$  during oxidation. The sediment-accumulation curves have small slopes, the original color of the oil changes insignificantly, and in most cases the corrosive aggressiveness of the oil with respect to copper and the contact resistance of the copper plates are reduced substantially. In contrast to inhibited oxidation, the kinetic curves reflecting the change in quantity of acids are smooth in the case of oils with anthranilic acid, with the amount of acids decreasing during oxidation. The latter is associated with depletion of the additive, which is acidic in nature.

During long-term aging of the oil, the passivating film formed on the copper surface disintegrates as the passivator is used up and oxidation products act, and further oxidation of the oil proceeds as though there were no additive present in it. This effect is particularly distinct in oxidation of oils with relatively small natural-inhibitor contents. To protect the adsorbed layer from attack by oxidation products and preserve it for a long time, an antioxidant additive is introduced into

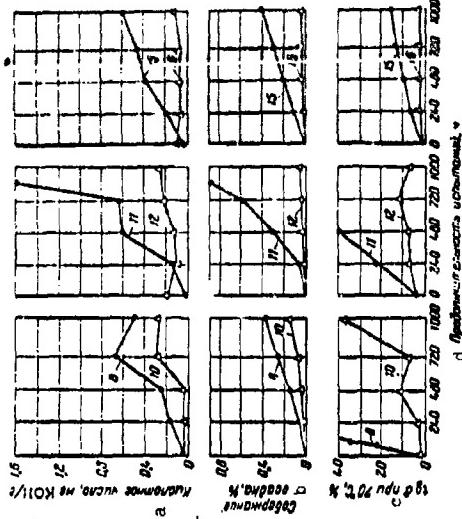


Fig. 3. Kinetic curves of indicator change during oxidation (1000 h) of transformer oils with additive mixes. (a) oil from Emba crudes without additives; (b) same with 0.02 [sic] anthranilic acid and 0.05% diisalicylidene-ethylenediamine (see Fig. 1 for other identifications).  
KEY: (a) acid number, mg of KOH/g; (b) sediment content, %; (c) tan  $\delta$  at 70°C, %; (d) test time, hours.

It was shown for the example of oils made from Aracataca and Buzhnominsk crudes (FIG. 3) that oxidation of the oil is inhibited sharply by use of a mixture consisting of 0.005% ionol, 0.1% phenylnaphthylamine.

It was established by the foregoing experiments that the effect is not observed when each of these additives is used alone. When the two additives are used the tangent of dielectric angle is stable even when separate introduction of 0.7% ionol or 0.05% phenylnaphthylamine does not produce the required effect.

Thus, when oil from Tuymazy crude is oxidized with each of the above additives,  $\tan \delta$  has already increased by 100% after 240-480 hours. On the other hand, when a mix consisting of 0.02% disalicylideneethylenediamine and 0.2% Ionol is introduced into the oil, its  $\tan \delta$  has increased only 43% after 1000 h of oxidation.

The stability of oil made from Emba crudes is improved substantially on addition of 0.05% disalicylideneethylenediamine and 0.02% anthranilic acid — something that could not be accomplished by use of a single additive. In this case, the action of the deactivator on the homogeneous catalyst — the dissolved copper — and that of the passivator on the heterogeneous catalyst — the metallic copper — are manifested simultaneously. Since there is an amino group in the anthranilic acid molecule, this passivator also has a weak inhibiting effect.

It should be noted that when the oil contains relatively large amounts of natural inhibitors, stability is improved only insignificantly by introducing an inhibiting additive together with the passivator (anthranilic acid).

Earlier experiments conducted with a hydrorefined oil made from Tuymazy crude confirm this. The results of laboratory tests to evaluate the effectiveness of additive combinations were confirmed in the bench tests with small transformers.

Additive mixes consisting of 0.05% anthranilic acid with 0.2% phenylnaphthylamine and 0.02% anthranilic acid with 0.05% disalicylideneethylenediamine improve the stability of oils made from Anasas'ev and Emba crudes. In particular, corrosion of copper is reduced substantially and destruction of specimens of solid insulating materials placed in the oil is retarded by a factor of 2-3.

A mix consisting of 0.05% anthranilic acid and 0.2% Ionol improves the stability of phenol-refined Tuymazy oil more effectively than a composition consisting of 0.05% disalicylideneethylenediamine and 0.2% Ionol. The difference in the effects of the two mixes is most noticeable in the relative amount of solid-insulation wear.

#### CONCLUSIONS

It has been shown that use of a passivating additive, anthranilic acid (0.05% by weight), in pure form and especially in mixes with antioxidant additives (Ionol, phenyl-naphthylamine) ensures better stability (especially electrophysical) of commercial transformer oils of various origins and chemical compositions than when only the inhibiting additive is used.

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Manu- script page	Symbol	Symbol	English Equivalent
102	A	A	aromatics
102	H	N	naphthenics

**INVESTIGATION OF ADDITIVES AND ADDITIVE  
OILS BY A POTENTIOMETRIC METHOD**

V.S. Luneva and L.N. Burderyuk

The advantages of using a potentiometric method to evaluate the acid-base properties of additives and oils containing additives and their wearability under the operating conditions of various types of engines have been confirmed by a number of research studies [1, 2].

**ALKALINITY OF ADDITIVES**

The alkalinity of an additive is governed by the presence of free and bound alkaline components [3, 4]. Additive alkalinities were determined by potentiometric titration using ammonia and calomel electrodes; 1 g. of additive dissolved in 40 ml. of an alcohol-benzol mixture (1:2) was decomposed by a 0.1N alcoholic solution of HCl. The titration curves appear in Figs. 1-5. As we see, the nature of the curves changes in accordance with the type of the salts decomposed: alkylsalicylate (ASK, NiASK), alkylphenol (VNIIM NP-339, VNIIM NP-370p), sulfonate (PMS'Ya), thiophosphate (DF-11, Santolube 493), etc. The length of the horizontal segment on the titration curve is proportional to the concentration of the salts decomposed during titration of equal weights. Additives containing the cations Ca, Ba, and Fe have an alkaline reaction ( $pE = 8-13$  or  $\Delta pH = 1-6$ ).

Alkylsalicylate (ASK, NiASK) and sulfonate (PMS'Ya) additives are characterized by free alkalinity (see initial segments of curves in Figs. 1-6). The alkylphenol additives (VNIIM NP-339, VNIIM NP-370p, etc.) do not contain free alkalies.

The titration curves indicate that most of the additives are totally decomposed at electromotive forces [emf] (EDC) of about 100-120 mV (pH = 2).

It is known that the total alkalinity [T.A.] (0.1N.) of most additives is equivalent to the content of metal present in them. This is confirmed by the data given in Table 1.

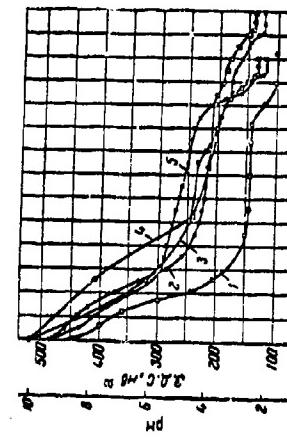


FIG. 1. Potentiometric titration curves of MASK additive. 1-5) various batches.  
KEY: (a) emf, mV; (b) volume of 0.1N HCl used in titration, ml.

However, this equivalence is not observed in the PMS'Ya and VNII NP-370 additives, so that alkalinity should generally be used to determine the effective content of metal, i.e., the content equivalent to that part of the additive that participates in neutralization of fuel combustion products. In introducing additive mixtures into the oil, it must be remembered that the sulfate ash content of PMS'Ya additive exceeds its total alkalinity by a factor of almost 1.5. PMS'Ya additives have different alkalinities at a given ash content (see Table 1). The increase in the ash content of the oil as the additives are depleted is one of the causes of increased wear, while inadequate alkalinity is among the causes of increased sludging. Hence it is necessary to check alkalinity and ash content in both the original additive oils and oils in use in engines.

#### INVESTIGATION OF ADDITIVES OF THE ZINC DITHIOPHOSPHATE TYPE

Owing to the aphoteric properties of zinc, its salts are decomposed by both acids and alkalies. Figure 4 presents potentiometric titration curves of zinc dialkyl- and diaryldithiophosphates, showing that the potential jump in acid titration

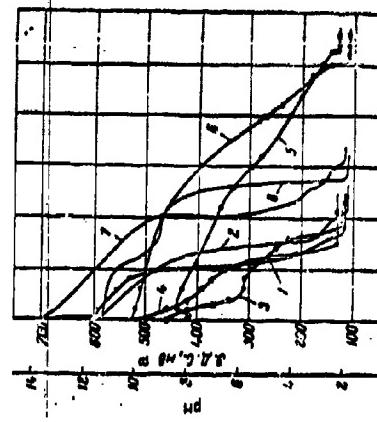


FIG. 2. Potentiometric titration curves of a-alkylphenol additives. 1-2) TRIATIN-319; 3,4) VNII NP-370; 5,6) VNII NP-370p; 7) VNII NP-380; 8) APB.  
KEY: (a) emf, mV; (b) volume of 0.1N HCl used in titration, ml.

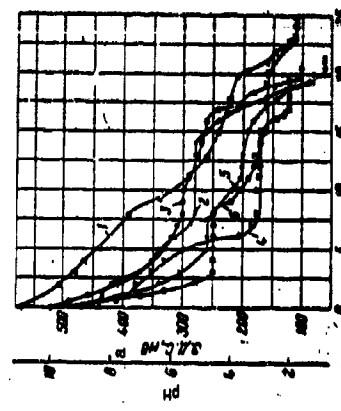


FIG. 3. Potentiometric titration curves of PMS'Ya additives. 1-6) various batches.  
KEY: (a) emf, mV; (b) volume of 0.1N HCl used in titration, ml.

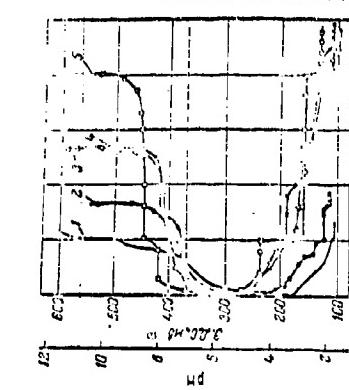


FIG. 4. Curves of potentiometric titration of zinc thiophosphate with alkali and acid:  
1) VNIIT-339; 2) DS-20A;  
3) Tsiatin-339; 4) Santolube  
1360; 5) Ecolube 99.  
KEY: (a) emf, mV; (b) volume  
of KOH or HCl added in  
titration, ml.

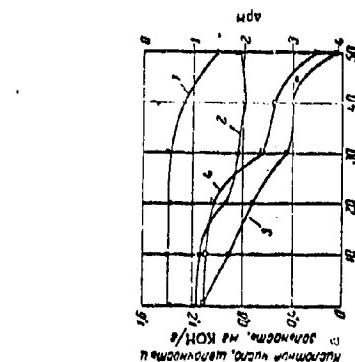


FIG. 5. Wearability of AS-9.5 oil with G-16 Orobis-Jloa 267 on IRG-2 bench installation.  
1) sulfate ash content; 2)  
total acid number; 3) total  
alkalinity; 4) reaction.  
KEY: (a) acid number, alka-  
linity, and ash content, mg  
of KOH/g; (b) time, hours.

Table 1. Results of Potentiometric Determination of Reaction, Alkalinity, Effective Metal and Sulfate-Ash Contents in Additives

a. Product	Components			base weight	P <sub>2O<sub>5</sub></sub> , %	A <sub>6</sub> , %
	b	c	d			
AASH-30	.....	3.2	6.28	7.00	62.1	0.1
ACK-70	.....	3.8	6.15	8.20	62.0	0.0
MACK	.....	1.6	1.2	9.15	58.3	0.1
WUCH	.....	2.3	1.58	11.5	53.5	0.1
E. HUZH	.....	1.0	1.11	11.5	53.5	0.1
R. BIRIN III-350 <sup>a</sup>	.....	4.5	1.65	12.7	55.0	0.1
R. BIRIN III-350 <sup>b</sup>	.....	0.8	1.90	12.7	55.0	0.1
R. BIRIN III-350 <sup>c</sup>	.....	2.7	1.55	16.9	58.9	0.1
C. BIRIN III-50	.....	3.5	1.91	1.11	52.0	0.1
F. RULATIN-339	.....	3.2	1.75	5.09	41.2	0.1
F. RULATIN-339	.....	4.2	1.65	5.05	57.0	0.1
AOB	.....	6.3	7.35	9.54	52.3	0.1
MACK	.....	1.3	0.51	17.6	51.0	0.1

\*Determined by turning the additive with sulfuric acid.

\*\*Barium.

\*\*\*Magnesium.

KEY: (a) additive; (b) reaction, +pH; (c) alkalinity,  
% of KOH/g; (d) content, %; (e) calcium  
ash; (f) sulfate ash; (g) difference of KOH/g;  
ash (B); (h) ASH-6v; (i) ASH-7v; (j) MACK;  
(k) PMS-Ya; (n) VNIIT MP-370; (o) VNIIT NP-310; (p)  
Tsiatin-339; (q) AFB; (r) MSZh.

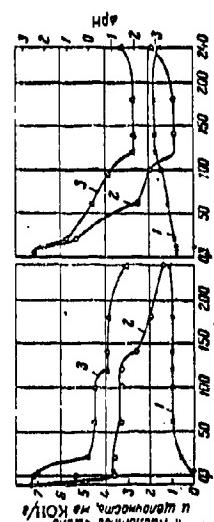


FIG. 6. Wearability of M14-V oil (DS-14 with  
3% Tsiatin-339, 2% PMS-Ya, and 0.03% PMS-20A)  
in SMP-14 engine during cycles I and IV. 1) total  
acid number; 2) total alkalinity; 3) reaction.  
KEY: (a) acid number and alkalinity, mg of KOH/g;  
(b) cycle I, hours; (c) cycle IV, hours.



The potential jump is sharper on decomposition of the additives by alkali according to the reaction



Two potential jumps can be brought out in titration of small weights (0.1g) (see FIG. 4); they correspond to decomposition of the salts with formation of thiophosphoric acid and potassium thiophosphate or decomposition of salts with various alkyl radicals.

Total decomposition of zinc dithiophosphates by alkali is accompanied by a potential jump in the 460-500 mV range (pH = 9-12). The total acid number [T.A.N.] (0.K.4.) of the additive is determined from the amount of alkali used in titrating to 520 mV (pH = 10).

The total acid number is converted by the formulas given below to the zinc and phosphorus contents.

$$0.K.4 = \frac{V_1 - V_2}{1} \text{ mg of KOH/g}$$

where  $V_1$  is the amount of 0.1N alcoholic KOH solution used in titrating the additive solution to pH = 10 (520 mV) in ml;  $V_2$  is the amount of the same solution used in titrating the solvent (blank experiment) to pH = 10 (520 mV), in ml;  $T$  is the titer of the 0.1N KOH solution, in g/ml; 1 is the weight of the additive sample, g.

The zinc content is

$$x_1 = \frac{0. K. 4 \cdot 0.583}{10} \%$$

The phosphorus content is

$$x_2 = \frac{0. K. 4 \cdot 0.655}{10} \%$$

where 0.583 is a coefficient obtained by dividing the atomic weight of zinc by the molecular weight of KOH; 0.55 is a coefficient obtained by dividing the atomic weight of phosphorus by the molecular weight of KOH; 10 is the coefficient for conversion to percent.

Example. For the Sanzilite 493 additive:

$$0.K.4 = \frac{(293-6) \cdot 1.19}{1} = 151.5 \text{ mg of KOH/g}$$

$$x_1 = \frac{151.5 \cdot 0.583}{56.1 \cdot 10} = \frac{151.5 \cdot 0.583}{561} = 8.88\% \\ x_2 = \frac{151.5 \cdot 0.655}{56.1 \cdot 10} = \frac{151.5 \cdot 0.655}{561} = 8.33\%$$

The advantages of the potentiometric method of determining zinc and phosphorus in the additives over the spectral, gravimetric, and calorimetric methods [7, 8] include the simplicity and speed of the analysis and the absence of burning or oxidation and precipitation losses; the time for a determination (1.5-2 hours) is smaller by a factor of more than 20 than in other presently known methods [7, 8, 9].

In Table 2, the results of potentiometric determination of zinc and phosphorus in the additives are compared with the

Table 2. Results of Quantitative Determination of Phosphorus and Zinc in Additives by Various Methods

a. additive	b. conversion of zinc and phosphorus to metal			c. conversion of zinc and phosphorus to phosphate		
	c	d	e	f	g	h
Gamma-493	1.12	1.12	1.09	1.09	82	82
Gamma-0-355	0.44	0.45	0.46	0.46	82	82
Gamma-100	1.12	1.12	1.09	1.09	82	82
Gamma-11	0.51	0.52	0.53	0.53	82	82
Gamma-HM-36	2.13	2.13	2.03	2.03	82	82
TAJAHN	2.16	2.09	2.07	2.07	82	82
Eltac, organic additive 60% zinc and phosphorus quinone	0.16	0.15-0.18	0.17	0.17	105-118	105-118
DTPA					82	82
DP-LL					82	82
NP-354					82	82
VNIIT NP-354					82	82
oil containing test specimen of zinc dithiophosphate					82	82

\*Determined calorimetrically (GOST [All-Union State Standard] 9827-61).

KEY: (a) specimen; (b) content, %; (c) potentiometric method; (d) spectral method (GOST 9466-63); (e) spectrophotometric or gravimetric method (GOST 1899-61); (f) Santicube h93; (h) Orobis-010a-267; (i) Lubrisol 1300; (j) DP-LL; (k) VNIIT NP-354; (m) oil containing test specimen of zinc dithiophosphate.

results obtained by other methods. Table 3 presents the results of potentiometric determination of reaction, total acid number, and phosphorus and zinc content. It shows that all of the thiophosphoric additives analyzed have an acid reaction due to the presence of the strong thiophosphate anion. The phosphorus content of the imported additives is considerably higher than that in the Soviet additives. The error of the results of determination varies: + 0.02% for the foreign additives and 0.9, + 0.5% for the domestic ones; the spread of the determination ranges from 2.4 to 9%. The potentiometric method can be used to establish the presence of small amounts of zinc and phosphorus (tenths and hundredths of a percent).

The presence of sulfur in these additives does not interfere with the potentiometric determination of zinc and phosphorus; here the emf values measured with the antimony electrode are not distorted.

In the whole, the above features of recommending all-additive potentiometric titration of zinc dithiophosphate for quantitative determination of phosphorus and zinc in the additives. This method can be used in research, standardization, and to control additive quality.

Potentiometric investigation of various types of additives enabled us to compute coefficients for conversion from total alkalinity to effective content of metal and sulfate at, as

Table 3. Results of Determination of Reaction, Total Acid Number, and Phosphorus and Zinc Contents in Metal-Dithiophosphate-Type Additives by Potentiometric Method

a additive	b	c	d	e	f
	Hydros.	Precip.	Oxide and KOH/g	content, %	as
f Cameline-408	1.01	-1.6	151.5	8.33	8.31
6 Opoider-Oloa-267	1.04	-1.1	152.7	8.31	8.36
h Meforane-1360	1.04	-1.2	153.2	8.42	8.39
1 Santolite-359	1.04	-2.1	159	7.45	8.09
1.2 Dithiophosphate of zinc	1.04	-0.8	159	7.45	8.09
1.3 Dithiophosphate of zinc	0.91	-1.9	217.3	11.98	12.59
1.4 Dithiophosphate of zinc	1.04	-1.8	217.9	11.98	12.59
1.5 Dithiophosphate of zinc	1.04	-2.2	85.2	3.76	3.87
1.6 Dithiophosphate of zinc	0.91	-1.7	98.9	4.96	5.21
1.7 Dithiophosphate of zinc	0.91	-3.7	105	5.82	6.18
1.8 Dithiophosphate of zinc	0.91	-3.7	45.5	2.75	2.87
1.9 Dithiophosphate of zinc	1	-3.3	1.38	0.07	0.08
2.0 Dithiophosphate of zinc	1	-1.9	2.90	0.05	0.17

KEY: (a) specimen; (b) weight taken, g; (c) reaction; (d) total acid number, mg of KOH/g; (e) content, %; (f) Santolite-493; (g) Orobis-Oloa-267; (h) Lubritisol-1360; (i) Ecolube-95; (j) LANI; (k) DF-11; (l) specimen ...; (m) VNII NP-354; (n) oil containing experimental specimen of zinc dithiophosphate.

Table 4. Conversion Factors K

a additive	b	c	Oxide and KOH/g						h
			1	2	3	4	5	6	
LACK, MACK, INCK, SHINN HILL-30, SHINN HILL-30, AOS	1.04	1.21	250	0.334	0.333	—	—	—	—
k SHINN HILL-30, AOS	1.04	1.22	2.08	0.519	0.518	1.02	—	—	—
m SHINN HILL-30, AOS	0.91	0.17	4.57	0.32	0.32	4.55	—	—	—
n Cameline-408, Oloa- 95, Santolite-493, DF-11	0.91	0.14	1.71	0.005	0.005	2.47	0.53	—	0.35

Working formula:  $M(MeSO_4) = T.A \cdot K \cdot 10$ , % (here Me is the metal content and  $Mes_4$  is the sulfate-ash content). Working formula:  $T.A = M(MeSO_4) \cdot K \cdot 10$ , mg of KOH/g. Working formula:  $Zn(P) = (M_A \cdot N \cdot K)^{1/2}$ , %.  
KEY: (a) additive; (b) element determined; (c) from total alkalinity to content, %; [cont'd on page 118]

[Key to Table 4, cont'd.] (1) of metals; (e) of sulfide ash; (f) from metal content; (g) and sulfate-ash content; (h) total alkalinity; (i) from alkalinities of (1) and back (2); (j) from total acidity to content of (1); (k) ASK, MSK, PMSV, VNII NP-370P; (l) calcium; (m) VNII NP-370, TSIAZIM-339, APM-1; (n) barium; (o) Ecolube-856, NSZPK [synthetic fatty acids]; (p) magnesium; (q) Santolite-493, Cetac-267, D-211; (r) zinc; (s) phosphorus.

#### ON THE QUESTION OF OIL SERVICE LIFE IN ENGINES

The wearability of oils with additives is the basic question to be resolved in establishing oil-change intervals for engines, and many investigators are working on it [20-13].

At one time [1], the question was raised as to the use of the potentiometric method to rate the acid-base properties of additive oils and their wearability in various types of engines.

The additive consumption computed from the change in oil ash content does not agree with the actual consumption, as shown in Figs. 5 and 7 and in Table 4.

As we see, alkalinity does not vary in proportion to additive content during depletion in engines. Alkalinity varies as a function of additive type, and, as a rule, the reaction (aph) changes from alkaline to acid or almost neutral (see Figs. 5-7).

Then, as we know, the viscosity of the oil and its coking capacity increase (Fig. 7b).

When the engines are operated under the same conditions (for example, the GAZ-51), equal quantities of the alkaline components are depleted, despite the difference in compositions (see Table 5). The amounts of sludge and wear are somewhat smaller at higher oil alkalinity.

Without additives, DS-14 oil services the YAK-204 engine for no more than 20 hours, and with additives for 85 hours [1]. Sludging appears at alkalinity not exceeding 0.4 mg of KOH/g. The alkalinites of the original oils were not the same (the oil used in the DS-14 engine was more alkaline).

DS-14 oil with 3% TSIAZIM-339, 2% PMSV, and 0.005% PMS-200A provides for normal operation of the DS-14 engine for 300 hours (see Fig. 6). Sludge appears after 500 hours in the Kolomna Plant's D-211 engine, with alkalinity not exceeding 0.4 mg of KOH/g. The alkalinites of the original oils were not the same (the oil used in the DS-14 engine was more alkaline).

JIL M-20F (MS-20 with 2.5% TSIAZIM-339, 1.5% PMSV, and 0.005% PMS-200A) works for 1360 hours in the M-7 engine [2].

Table 5. Physicochemical and Motor Properties of Additive Oil Before and After Use in GAZ-51 Engine

**KEY:** (a) oil with additives; (b) acid-base properties; (c) total alkalinity, mg of KOH/g; (d) total acid number, mg of KOH/g; (e) wearability; (f) motor rating; (g) amount of carbon on rings and piston, %; (h) cylinder wear (by method of lines),  $\mu$ ; (i) weight loss of first piston ring, g; (j) oil burner, g/h; (k) viscosity at 100°C, cSt; (l) content, %; (m) coke, %; (n) ash; (o) M-100B (AS-9.5 + 0.7% Santon-493; (p) original; (q) after 600 hours of use; (r) M-100B (AS-9.5 + 3.0% ASK + 1.2% DP-1 + 0.0035 PMS-200A); (s) M-100A (AS-0.5 + 1.5% ASK + 1.2% DP-1 + 0.0033 PMS-200A).

stages, each lasting 50 hours) instead of the specified 100 hours. The engine was in satisfactory condition.

We see from the data obtained (FIG. 5-7) that the alkalinity decrease depends not only on the duration of engine running, but also on running conditions.

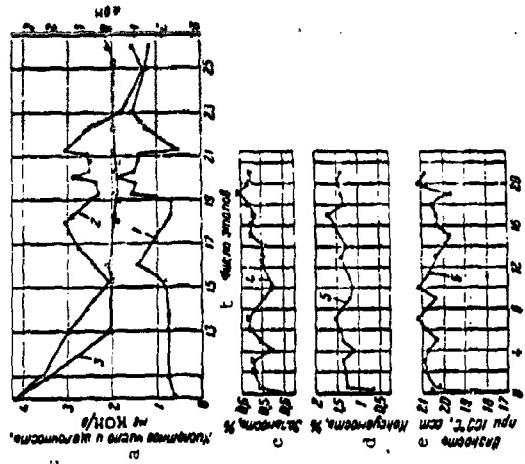


Fig. 7. Wearability of M-20Y oil (M-20 with 2.5% TISATIN-M-339, 15% PMS-Ya and 0.005% PMS-20A) in M-20 engine. 1) total acid numbers; 2) total alkalinity; 3) reaction; 4) ash; 5) coding number; 6) viscosity of KOH/B; (b) number of stages of KOH/B; (c) ash; (d) coding capacity; (e) viscosity at 100°C, cst.

סימן י

1. The possibility and expediency of direct titration of additives to 120-100 mV ( $\text{pH} = 2$ ) to determine total alkalinity and their effective metal and sulfate-ash content; (the latter for those additives in which alkalinity is equivalent to ash) were demonstrated.
  2. The sulfonate additives (PMSYa) differ sharply from one another in their effectiveness, hence, in successive cal-

compounds, ranging up to 9% (converted to calcium sulfate).

3. The zinc thiophosphates that were analyzed had an acid reaction; their acid numbers were proportional to phosphorus and zinc content; the imported additives contained substantially more phosphorus and zinc than the Soviet ones. A fundamentally new method was proposed for quantitative determination of phosphorus and zinc by titrating zinc dithiophosphate to the potential jump or, in the general case, to 520 mV (pH = 10). The error of the method is 0.02 to 0.25%, its reproducibility from +2.4 to +9%, and the time required for the determinations 1.5-2 hours.

4. A study of the acid-base properties of additive oils in various types of engines brought out a disproportionality between the depletion of the alkaline component and the accumulation of acidic compounds; alkalinity does not vary in proportion to ash content. It was established that in engines operated under uniform conditions (GAZ-51) but on oils with different additive mixes, the alkalinity reserve varies in about the same way. A relation was detected between alkalinity change and the cleanliness and wear of certain engines.

5. The expediency of using the potentiometric method of wearability rating for additive oils to arrive at change intervals for these oils in various types of engines was demonstrated.

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